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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 03:28:19 ; Search time 222.28 Seconds

(without alignments)
347.126 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
Sequence: 1 atgaaaaaattatttcaaa.....gtccaatgaagcaagtgca 90

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 930621 segs, 428662619 residues

Word size: 0

Total number of hits satisfying chosen parameters: 989696

Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

N.Geneseq_1101:*

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2: /SIDS2/gcgdata/geneseq/geneseq/NA1981.DAT:*

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5: /SIDS2/gcgdata/geneseq/geneseq/NA1984.DAT:*

6: /SIDS2/gcgdata/geneseq/geneseq/NA1985.DAT:*

7: /SIDS2/gcgdata/geneseq/geneseq/NA1986.DAT:*

8: /SIDS2/gcgdata/geneseq/geneseq/NA1987.DAT:*

9: /SIDS2/gcgdata/geneseq/geneseq/NA1988.DAT:*

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11: /SIDS2/gcgdata/geneseq/geneseq/NA1990.DAT:*

12: /SIDS2/gcgdata/geneseq/geneseq/NA1991.DAT:*

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14: /SIDS2/gcgdata/geneseq/geneseq/NA1993.DAT:*

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16: /SIDS2/gcgdata/geneseq/geneseq/NA1995.DAT:*

17: /SIDS2/gcgdata/geneseq/geneseq/NA1996.DAT:*

18: /SIDS2/gcgdata/geneseq/geneseq/NA1997.DAT:*

19: /SIDS2/gcgdata/geneseq/geneseq/NA1998.DAT:*

20: /SIDS2/gcgdata/geneseq/geneseq/NA1999.DAT:*

21: /SIDS2/gcgdata/geneseq/geneseq/NA2000.DAT:*

22: /SIDS2/gcgdata/geneseq/geneseq/NA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	14	15.6	20	AA205048	PCR primer used to
2	14	15.6	26	AAV07952	Helicobacter pylori
3	14	15.6	26	AAV07922	Helicobacter pylori
4	14	15.6	27	AAV07937	Helicobacter pylori
5	14	15.6	47	AA268842	Human map-related
6	13	14.4	17	AA092084	Human reniformis
7	13	14.4	20	AA353307	PCR primer used to
8	13	14.4	23	AAH27610	Human lipoprotein
9	13	14.4	24	AAH55939	Human SCNA PCR-SS
10	13	14.4	26	AA068537	B. thuringiensis 33
11	13	14.4	26	AA067312	Alzheimer's disease

12	13	14.4	26	AA067342	Alzheimer's disease
13	13	14.4	31	AAV07957	Helicobacter pylori
14	13	14.4	34	AA153377	AZF A5F cosmid c10
15	13	14.4	35	AA061207	Coxsackie virus B
16	13	14.4	36	AA090560	Dissociation trans
17	13	14.4	38	AA134130	I3L promoter-HIV-
18	13	14.4	39	AA035630	HIV-2 env 3' fragm
19	13	14.4	39	AA035353	PCR primer HIV2B2
20	13	14.4	39	AA099138	Plasmid pomptc PC
21	13	14.4	40	AA048717	Insecticidal prote
22	13	14.4	40	AA261322	Primer 3A used to
23	13	14.4	40	AA261323	Primer 3B used to
24	13	14.4	41	AAV51009	Maize polymorphic
25	13	14.4	41	AAV51011	Maize polymorphic
26	13	14.4	50	AA125074	Human gene signatu
27	13	14.4	50	AA052184	Synthetic plasmid
28	13	14.4	50	AA052048	Synthetic plasmid
29	13	14.4	17	AA070046	Human fil1 VEGF re
30	12	13.3	17	AA070047	Human fil1 VEGF re
31	12	13.3	17	AA021453	Integrin alpha 6 s
32	12	13.3	17	AA021454	Integrin alpha 6 s
33	12	13.3	17	AA021455	Integrin alpha 6 s
34	12	13.3	17	AA030888	Hammerhead ribozym
35	12	13.3	17	AA030889	Hammerhead ribozym
36	12	13.3	17	AA030890	Hammerhead ribozym
37	12	13.3	17	AA030891	Hammerhead ribozym
38	12	13.3	18	AA089921	Human survivin DNA
39	12	13.3	19	AA062806	Env gene 5' primer
40	12	13.3	19	AA069625	Human biallelic ma
41	12	13.3	19	AA275126	HIV RNA translati
42	12	13.3	20	AA005907	Staphylococcus aur
43	12	13.3	20	AA080813	Corn kernel oil co
44	12	13.3	20	AAV72697	PCR primer used to
45	12	13.3	20	AA032327	

ALIGNMENTS

RESULT 1	
AA205048	AA205048 standard; DNA; 20 BP.
XX	XX
XX	AA205048;
XX	07-OCT-1999 (first entry)
XX	PCR primer used to amplify an ORF of Chlamydia trachomatis.
XX	
XX	Vaccine, eye disease; conventional trachoma; nonendemic trachoma;
XX	paratrachoma; inclusion conjunctivitis; genital disease; perinephritis;
XX	nongonococcal urethritis; epididymitis; cervicitis; salpingitis; PCR primer;
XX	bartholinitis; pneumonia; venereal lymphogranulomatosis; ss.
XX	
XX	Synthetic.
XX	Chlamydia trachomatis.
XX	
XX	W09928475-A2.
XX	
XX	10-JUN-1999.
XX	
XX	27-NOV-1998; 98WO-IB01939.
XX	
XX	04-NOV-1998; 98US-0107077.
XX	28-NOV-1997; 97FR-0015041.
XX	17-DEC-1997; 97FR-0016034.
XX	
XX	(GEST) GENSET.
XX	
XX	Griffais R;
XX	
XX	WPI, 1999-371125/31.
XX	

PT Genome sequence of Chlamydia trachomatis
XX
PS Disclosure; Page 1738; 1755pp; English.
XX
CC PCR primers AA201426-206209 were used to amplify open reading frames
CC (ORFs) of the genome of Chlamydia trachomatis (see AA201425). These ORFs
CC encode polypeptides (see AA36754-Y37949) which can be used as vaccines
CC against Chlamydia trachomatis. Antisense and ribozyme sequences
CC can also be used to control growth of the microorganism. Chlamydia
CC trachomatis is responsible for a large number of diseases, e.g. eye
CC diseases such as conventional trachoma, nonendemic trachoma,
CC paratrachoma, and inclusion conjunctivitis; genital diseases such as
CC nongonococcal urethritis, epididymitis, cervicitis, salpingitis,
CC perihepatitis, Bartholinitis; pneumopathy in breast feeding infants;
CC and venereal lymphogranulomatosis. The polypeptides of the
CC invention may be of use in treating these diseases.
XX
SQ Sequence 20 BP; 5 A; 5 C; 5 G; 5 T; 0 other;

Query Match 15.6%; Score 14; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 9.4e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 54 tattgttgagcaaa 67
|||||
Db 4 tattgttgagcaaa 17

RESULT 2
AAV07952/C
ID AAV07952 standard; DNA; 26 BP.
XX
AC AAV07952;
XX
DT 02-FEB-1999 (first entry)
XX
DE Helicobacter pylori polypeptide GHPO 1414 5' DNA primer.
XX
DE GHPO 1414: infection; gastritis; ulcer; vaccine; diagnosis;
XX therapy; PCR; primer; ss.
XX
KM Synthetic.
XX
OS Helicobacter pylori.
XX
OS WO9843479-A1.
XX
PN 08-OCT-1998.
XX
PD 31-MAR-1998; 98WO-US06421.
XX
PF 01-APR-1997; 97US-0834666.
XX
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX (INNR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX
PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX WPI: 1998-568251/48.
XX
XX New isolated Helicobacter polynucleotides - used to develop products
XX for the diagnosis, prevention and treatment of Helicobacter
XX infections and gastroduodenal diseases
XX
XX Claim 5; Page 145; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07954) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV07921) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73032) designated GHPO 1414.
CC The isolated polynucleotide, and encoded polypeptide, can be used
CC to develop vaccines for the treatment and prevention of Helicobacter

CC Infections.
XX
SQ Sequence 26 BP; 14 A; 5 C; 4 G; 3 T; 0 other;

Query Match 15.6%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 9.3e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 40 ttctcatgtttct 53
|||||
Db 18 TTTTCATGTTTCT 5

RESULT 3
AAV07922/C
ID AAV07922 standard; DNA; 26 BP.
XX
AC AAV07922;
XX
DT 02-FEB-1999 (first entry)
XX
DE Helicobacter pylori polypeptide GHPO 386 5' DNA primer.
XX
DE GHPO 386: infection; gastritis; ulcer; vaccine; diagnosis; therapy;
XX PCR; primer; ss.
XX
KM Synthetic.
XX
OS Helicobacter pylori.
XX
OS WO9843479-A1.
XX
PN 08-OCT-1998.
XX
PD 31-MAR-1998; 98WO-US06421.
XX
PF 01-APR-1997; 97US-0834666.
XX
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX (INNR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX
PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX WPI: 1998-568251/48.
XX
XX New isolated Helicobacter polynucleotides - used to develop products
XX for the diagnosis, prevention and treatment of Helicobacter
XX infections and gastroduodenal diseases
XX
XX Claim 5; Page 137; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07924) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV72001
CC) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73032) designated GHPO 386. The
CC isolated polynucleotide, and encoded polypeptide, can be used to
CC develop vaccines for the treatment and prevention of Helicobacter
XX infections.
XX
SQ Sequence 26 BP; 15 A; 5 C; 4 G; 2 T; 0 other;

Query Match 15.6%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 9.3e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 40 ttctcatgtttct 53
|||||
Db 20 TTTTCATGTTTCT 7


```

RESULT 4
AAV07937/C
ID AAV07937 standard; DNA; 27 BP.
XX
AC AAV07937;
XX
DT 02-FEB-1999 (first entry)
XX
DE Helicobacter pylori polypeptide GHP0 896 5' DNA primer.
XX
KM GHP0 896; infection; gastritis; ulcer; vaccine; diagnosis;
XX therapy; PCR; primer; ss.
XX
OS Synthetic.
XX Helicobacter pylori.
XX
PM W09843479-A1.
XX
PD 08-OCT-1998.
XX
PF 31-MAR-1998; 98WO-US06421.
XX
PR 01-APR-1997; 97US-0834666;
XX 01-APR-1997; 97US-0831310.
XX
PA (HIMA-) HUMAN GENOME SCI INC.
XX (IMMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX
PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX
DR WPI; 1998-568251/48.
XX
PT New isolated Helicobacter polynucleotides - used to develop products
XX for the diagnosis, prevention and treatment of Helicobacter
XX infections and gastroduodenal diseases
XX
PS Claim 5; Page 141; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07939) in the PCR
XX amplification of Helicobacter, e.g. Helicobacter pylori, genomic
XX DNA in order to obtain DNA (see AAV07916) encoding the unprocessed
XX form of a 76 kDa polypeptide (see AAW73027) designated GHP0 896.
XX The isolated polynucleotide, and encoded polypeptide, can be used
XX to develop vaccines for the treatment and prevention of Helicobacter
XX infections.
XX
SQ Sequence 27 BP; 14 A; 5 C; 4 G; 4 T; 0 other;

Query Match 15.6%; Score 14; DB 19; Length 27;
Best Local Similarity 100.0%; Pred. No. 9.3e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 40 ttctcatgtttct 53
Db 18 TTTTCATCTTTCT 5

```

```

OS Homo sapiens.
XX
FH Key Location/Qualifiers
XX variation replace(24,G)
XX FT /*tag= a
XX FT /standard_name= "single nucleotide polymorphism"
XX
PM W09954500-A2.
XX
PD 28-OCT-1999.
XX
PF 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
XX 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
XX
DR WPI; 2000-013267/01.
XX
PT Novel biallelic markers used to construct a high density disequilibrium
XX map of the human genome
XX
PS Claim 3; Page 912; 2745pp; English.
XX
CC AA265654 to AA269578 represent human biallelic markers from the present
XX invention, which contain a polymorphic base at position 24 of their
XX nucleotide sequences. AA269579 to AA277440 represent amplification
XX primers for the biallelic markers. The biallelic markers of the
XX invention have a variety of uses: they can be used for high density
XX mapping of the human genome, and in complex association studies and
XX haplotyping studies which are useful in determining the genetic basis
XX for disease states. Compositions and methods of the invention can also
XX be useful for the identification of the targets for the development of
XX pharmaceutical agents and diagnostic methods, as well as the
XX characterisation of the differential efficacious responses to and side
XX effects from pharmaceutical agents acting on a disease as well as other
XX treatment.
XX N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
XX and 3367, are not actually given a sequence in the Sequence Listing
XX from the present invention.
XX
SQ Sequence 47 BP; 13 A; 13 C; 6 G; 15 T; 0 other;

Query Match 15.6%; Score 14; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 9.1e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 75 aatgaagcaagt 88
Db 43 AATGAAGCAAGTG 30

```

PD 23-MAY-1995.
XX
XX 29-DEC-1989; 89US-0458952.
XX
XX 29-DEC-1989; 89US-0458952.
PR 20-AUG-1992; 92US-0933017.
PR 17-JUN-1993; 93US-0079700.
PR 14-DEC-1993; 93US-0167650.
XX
PA (UYGE-) UNIV GEORGIA RES FOUND INC.
XX
PI Cormier MJ, Lorenz WJ;
XX
XX WPI; 1995-199740/26.
XX
PT New recombinant Renilla luciferase polypeptide - used as a
PT luminescent tag, partic in bio-luminescence assays and for the prodn
PT of antibodies
XX
PS Disclosure; Fig. 4; 18pp; English.
XX
XX This 17-mer oligonucleotide DNA probe, along with probe-2 (AAQ92085)
CC are used to screen an R. reniformis cDNA library to isolate cDNA
CC encoding Renilla luciferase. The luciferase was then expressed
CC using E. coli.
XX
SQ Sequence 17 BP; 6 A; 0 C; 2 G; 9 T; 0 other;

Query Match 14.4%; Score 13; DB 16; Length 17;
Best Local Similarity 100.0%; Pred. No. 2.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattattt 16
|||||
Db 15 AAAAAATTATTT 3

RESULT 7
AAK95307
ID AAK95307 standard; DNA; 20 BP.
XX
AC AAK95307;
XX
DT 13-SEP-1999 (first entry)
XX
DE PCR primer used to amplify an ORF of Chlamydia pneumoniae.
XX
KW Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;
KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;
KW vaccine; neutralising epitope; PCR primer; ss.
XX
OS Synthetic.
OS Chlamydia pneumoniae.
XX
PN WO9927105-A2.
XX
PD 03-JUN-1999.
XX
PF 20-NOV-1998; 98WO-IB01890.
XX
PR 04-NOV-1998; 98US-0107078.
PR 21-NOV-1997; 97FR-0014673.
XX
PA (GEST) GENSET.
XX
PI Griflais R;
XX
XX WPI; 1999-357842/30.
XX
PT Genome sequence of Chlamydia pneumoniae
PS Page 1737; Disclosure; 1912pp; English.

XX
XX AAK91991-x97517 represent PCR primers used to amplify open reading
CC frames and other nucleic acid sequences from the genome of
CC Chlamydia pneumoniae (see AAK91990). C. pneumoniae causes respiratory
CC disease such as pneumonia and bronchitis and is thought to be a
CC contributing factor in heart disease, sarcoidosis, sinusitis, purulent
CC otitis media, erythema nodosum or pharyngitis. The polypeptides encoded
CC by the open reading frames of the C. pneumoniae genome (see AAK94584-
CC AAK95879) can be used in immunogenic compositions as vaccines. Vectors
CC containing C. pneumoniae nucleotides sequences can also be used as
CC immunogenic compositions, especially where the vector directs the
CC expression of a neutralising epitope of C. pneumoniae.
XX
SQ Sequence 20 BP; 5 A; 5 C; 5 G; 5 T; 0 other;

Query Match 14.4%; Score 13; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 2.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagtcg 89
|||||
Db 8 tgaagcaagtcg 20

RESULT 8
AAH27610
ID AAH27610 standard; DNA; 23 BP.
XX
AC AAH27610;
XX
DT 29-AUG-2001 (first entry)
XX

DE Human lipoprotein 105 PCR primer 2.

XX
XX Human; lipoprotein 105; cytosolic; anti-HIV; antiinflammatory;
KW immunomodulatory; cytochrome c structural domain; cancer;
KW haemopathy; human immunodeficiency virus; HIV; immunological disease;
KW inflammation; PCR primer; ss.
XX

OS Homo sapiens.

XX
XX WO200140483-A1.
XX
PN 07-JUN-2001.
XX
PD 27-NOV-2000; 2000WO-CN00503.
XX

PF 29-NOV-1999; 99CN-0124136.

PR (BIOR-) BIOROAD GENE DEV LTD SHANGHAI.

PA Mao Y, Xie Y;

PI WPI; 2001-374839/39.

XX
XX Human lipoprotein 105 containing cytochrome c structural domain and
PT encoded polynucleotide, applicable in diagnosis and treatment of
PT malignant tumor, hemopathy, HIV infection, immunological diseases and
PT various inflammation
XX
XX Example 2; Page 12; 40pp; Chinese.

XX
XX The invention relates to an isolated polypeptide of human lipoprotein
CC 105 containing a cytochrome c structural domain. The polypeptide
CC comprises a 951 amino acid sequence given in the specification, or its
CC fragment, analogue or derivative. The polypeptide and encoded
CC polynucleotide are useful in the diagnosis and treatment of malignant
CC tumours, haemopathy, HIV infection, immunological diseases and various
CC inflammatory diseases. The present sequence is a primer which was
CC used to isolate the polynucleotide encoding the polypeptide of the
XX invention.

Sequence 23 BP; 5 A; 1 C; 2 G; 15 T; 0 other;

Query Match 14.4%: Score 13; DB 22; Length 23;

Best Local Similarity 100.0%; Pred. No. 2.8e+03; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

33 ttattatgtttca 45
|||||
1 ttattatgtttca 13

RESULT 9

AAH55939

ID AAH55939 standard; DNA; 24 BP.

AAH55939;

04-SEP-2001 (first entry)

Human SCN1A PCR-SSCP PCR primer SEQ ID NO:183.

Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification; diagnosis; mutation; chromosome 2q23-q31; neurological disorder; anticonvulsant; neuroprotective; PCR primer; ss.

Homo sapiens.
Synthetic.

WO200138564-A2.

31-MAY-2001.

24-NOV-2000; 2000WO-CA01404.

26-NOV-1999; 99US-0167623.

(UYWC-) UNIV MCGILL.

Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;

WPI; 2001-355945/37.

Determining a predisposition to epilepsy and/or development of epilepsy comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA variant, equivalent, or mutation which shows a linkage disequilibrium -

Example 3; Fig 2; 268pp; English.

The present invention describes a method (M1) of determining an individual's predisposition to epilepsy and/or development of epilepsy, as well as predicting the individual's response to medication. The method comprises determining the genotype of at least one gene selected from SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium channel genes located on chromosome 2. The idiopathic generalised epilepsy (IGE) gene is more specifically localised on chromosome 2q23-q31. Compounds identified as modulators of the biological activity of SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy or other neurological disorders. They have anticonvulsant and neuroprotective activities. AAH55763 to AAH56164 and AAH9674 to AAH96679 represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers, oligonucleotides and proteins given in the exemplification of the present invention.

Sequence 24 BP; 2 A; 5 C; 3 G; 14 T; 0 other;

Query Match 14.4%: Score 13; DB 22; Length 24;

Best Local Similarity 100.0%; Pred. No. 2.8e+03; Mismatches 0; Indels 0; Gaps 0;

Qy 34 ttatgtttttcat 46
|||||
Db 8 ttatgtttttcat 20

RESULT 10

AAO68537/C

ID AAO68537 standard; cDNA; 26 BP.

AAO68537;

13-FEB-1995 (first entry)

B. thuringiensis 33kD delta-endotoxin N-terminal probe.

Insecticidal protein; delta-endotoxin; crystal; Coleoptera; Lepidoptera; Bacillus thuringiensis; ss.

Synthetic.

WO9413785-A.

23-JUN-1994.

13-DEC-1993; 93WO-US12144.

15-DEC-1992; 92US-0991073.

(NOVO) NOVO-NORDISK ENTOTEC INC.

Adams LF, Liu C, Lufurrow PA, Thomas MD;

WPI; 1994-217865/26.

New Bacillus thuringiensis strains - which produce new delta-endotoxin cpds used for the control of Lepidopteran and Coleopteran insect pests.

Example 9; Page 26; 47pp; English.

The N-terminal sequence of a 33kD delta-endotoxin isolated from B. thuringiensis EMC0075 (NRRL B-21019) or EMC0076 (NRRL B-21020) was determined (see AAR59764). Based on this sequence a 26mer oligonucleotide was designed for use as a probe (AAO68537) for cloning the delta-endotoxin gene.

Sequence 26 BP; 13 A; 2 C; 2 G; 9 T; 0 other;

Query Match 14.4%: Score 13; DB 15; Length 26;
Best Local Similarity 100.0%; Pred. No. 2.8e+03; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 30 aattttatgttt 42
|||||
Db 16 AATTTTATGTTT 4

RESULT 11

AAO67312

ID AAO67312 standard; DNA; 26 BP.

AAO67312;

14-FEB-2001 (first entry)

Alzheimer's disease-linked mitochondrial SNP PCR primer #12.

Human; mitochondrial genome; single nucleotide polymorphism; SNP; Alzheimer's disease; mtDNA; PCR primer; ss.

Homo sapiens.

PN WO200063441-A2.
 XX
 PD 26-OCT-2000.
 XX
 PF 19-APR-2000; 2000WO-US10906.
 XX
 PR 20-APR-1999; 99US-0130447.
 PR 22-OCT-1999; 99US-0160901.
 XX
 PA (MITO-) MITOKOR.
 PI Herrnstadt C, Davis RE;
 XX
 DR WPI; 2000-672748/65.
 XX
 PT Diagnosing a subject at the risk for or having Alzheimer's disease
 PT comprises determining at least one single nucleotide polymorphism in
 PT mitochondrial DNA associated with the disease in the sample from the
 PT subject -
 XX
 PS Example 2; Page 34; 89pp; English.
 XX
 CC The present invention describes a novel method for determining the risk
 CC of or diagnosing Alzheimer's disease using single nucleotide
 CC polymorphisms (SNPs) present in an individual's mitochondrial DNA
 CC (mtDNA). In addition, the SNPs identified can be used to identify agents
 CC suitable for use in treating Alzheimer's disease. Sequences
 CC AAC67301-C67610 are PCR primers used to demonstrate the method of the
 CC invention.
 CC
 SQ Sequence 26 BP; 8 A; 2 C; 5 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 26;
 Best Local Similarity 100.0%; Pred. No. 2.8e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 37 atgtttcatgtt 49
 ||||||||||||
 Db 7 atgtttcatgtt 19

RESULT 12
 AAC67342
 ID AAC67342 standard; DNA; 26 BP.
 XX
 AC AAC67342;
 XX
 DT 14-FEB-2001 (first entry)
 XX
 DE Alzheimer's disease-linked mitochondrial SNP PCR primer #42.
 XX
 KW Human; mitochondrial genome; single nucleotide polymorphism; SNP.
 KW Alzheimer's disease; mtDNA; PCR primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200063441-A2.
 XX
 PD 26-OCT-2000.
 XX
 PF 19-APR-2000; 2000WO-US10906.
 XX
 PR 20-APR-1999; 99US-0130447.
 PR 22-OCT-1999; 99US-0160901.
 XX
 PA (MITO-) MITOKOR.
 PI Herrnstadt C, Davis RE;
 XX
 DR WPI; 2000-672748/65.
 XX
 PT Diagnosing a subject at the risk for or having Alzheimer's disease

PT comprises determining at least one single nucleotide polymorphism in
 PT mitochondrial DNA associated with the disease in the sample from the
 PT subject -
 XX
 PS Example 4; Page 38; 89pp; English.
 XX
 CC The present invention describes a novel method for determining the risk
 CC of or diagnosing Alzheimer's disease using single nucleotide
 CC polymorphisms (SNPs) present in an individual's mitochondrial DNA
 CC (mtDNA). In addition, the SNPs identified can be used to identify agents
 CC suitable for use in treating Alzheimer's disease. Sequences
 CC AAC67301-C67610 are PCR primers used to demonstrate the method of the
 CC invention.
 CC
 SQ Sequence 26 BP; 8 A; 2 C; 5 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 26;
 Best Local Similarity 100.0%; Pred. No. 2.8e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 37 atgtttcatgtt 49
 ||||||||||||
 Db 7 atgtttcatgtt 19

RESULT 13
 AAV07957/G
 ID AAV07957 standard; DNA; 31 BP.
 XX
 AC AAV07957;
 XX
 DT 02-FEB-1999 (first entry)
 XX
 DE Helicobacter pylori polypeptide GPO 386 5' DNA primer.
 DE GPO 386; infection; gastritis; ulcer; vaccine; diagnosis; therapy;
 KW PCR; primer; ss.
 KW
 OS Synthetic.
 OS Helicobacter pylori.
 OS
 PN WO9843479-A1.
 XX
 PD 08-OCT-1998.
 XX
 PF 31-MAR-1998; 98WO-US06421.
 XX
 PR 01-APR-1997; 97US-0834666.
 PR 01-APR-1997; 97US-0831310.
 XX
 PA (HOMA-) HUMAN GENOME SCI INC.
 PA (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
 PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
 PI WPI; 1998-568251/48.
 XX
 DR
 XX
 PT New isolated Helicobacter polynucleotides - used to develop products
 PT for the diagnosis, prevention and treatment of Helicobacter
 PT infections and gastroduodenal diseases
 XX
 PS Example 3.B; Page 63; 184pp; English.
 XX
 CC This 5' primer was used with a 3' primer (see AAV07958) in the PCR
 CC amplification of Helicobacter pylori strain ORV2001 genomic DNA in
 CC order to obtain DNA (see AAV72001
 CC) encoding a 76 kDa polypeptide (see
 CC AAV73022) designated GPO 386. The primer pair includes a 5' clamp
 CC and BamHI and XhoI restriction enzyme recognition sequences for
 CC cloning purposes. The PCR product was ligated into vector pET28a,
 CC and recombinant polypeptide was expressed as a histidine-tagged
 CC fusion protein in E. coli host cells. The polypeptide can be used

CC to develop vaccines for the treatment and prevention of Helicobacter
 CC infections.

XX Sequence 31 BP; 14 A; 4 C; 6 G; 7 T; 0 other;

XX SQ

Query Match 14.4%; Score 13; DB 19; Length 31;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 41 ttcatgttttct 53
 |||||
 Db 31 TTTCATGTTTCT 19

RESULT 14

AA15377
 ID AA15377 standard; DNA; 34 BP.

XX AC AA15377;

XX DT 20-MAR-1996 (first entry)

XX DE AZF A5F cosmid clone exon 9/intron 9 boundary.

XX KW Azospermia factor; AZF; male infertility; YRRM gene;

XX KM Y-chromosome; ss.

XX OS Homo sapiens.

XX Key

XX exon

XX intron

XX /tag= a

XX /number= 9

XX /note= "Represents bases 1093-1109 of human YRRM gene"

XX /tag= b

XX /number= 9

XX /note= "Constitutes 5' end of approx. 450 bp intron 9"

XX W09511300-A2.

XX PD 27-APR-1995.

XX XX 24-OCT-1994; 94WO-GB02344.

XX PR 07-JUL-1994; 94GB-0013760.

XX PR 22-OCT-1993; 93GB-0021857.

XX PA (MEDI-) MEDICAL RES COUNCIL.

XX PI Chandley AC, Cooke HJ, Hargreave TB, Kun M, Sharkey AM;

XX DR WPI: 1995-170221/22.

XX Nucleic acid encoding the human azospermia factor, and probes and

XX PT antibodies specific for the sequence and encoded polypeptide - may

XX PT be used in the clinical diagnosis of male infertility

XX PS Disclosure: Fig 3; 40pp; English.

XX CC The intron/exon boundaries of a human YRRM gene encoding

XX CC azospermia factor were detd. by comparison of the sequences

XX CC found in cosmid A5F DNA obtd. from a Y-chromosome specific cosmid

XX CC library and the CDNA clone MK5 (AA087655) obtd. from an adult human

XX CC testis library.

XX SQ Sequence 34 BP; 9 A; 3 C; 7 G; 15 T; 0 other;

Oy 24 tactgtattttt 36
 |||||
 Db 19 tactgtattttt 31

RESULT 15

AA15377
 ID AA15377 standard; DNA; 35 BP.

XX AC AA15377;

XX DT 25-MAY-2001 (first entry)

XX DE Cocksackie virus B serotype B3 associated PCR primer SEQ ID 4.

XX KW Gene therapy; infectious virion; cardiactive; cardiac muscle; trophic;

XX KM cardiac disease; cardiac myocyte; PCR primer; ss.

XX OS Cocksackievirus.

XX DE19939095-A1.

XX PD 22-FEB-2001.

XX PF 18-AUG-1999; 99DE-1039095.

XX PR 18-AUG-1999; 99DE-1039095.

XX PA (UYTU-) UNIV TUEBINGEN EBERHARD-KARLS.

XX PI Kuepper J, Meyer R, Meyer-Ficca M, Kandolf R;

XX WPI: 2001-227548/24.

XX Recombinant RNA comprising heterologous gene in Cocksackie viral genome,

XX PT useful in gene therapy, specifically for targeting of cardiac myocytes

XX Example 3; Column 10; 12pp; German.

XX This invention describes a novel recombinant RNA molecule (I), at least

XX partly translatable in a target cell which comprises: (a) the

XX non-infectious genome (A) of Group B Cocksackie virus (CVB), particularly

XX serotype B3; and (b) at least one foreign gene (II) that can be developed

XX for a selected function in the target cell, e.g. for gene therapy. The

XX invention also describes (1) a recombinant infectious virions (V),

XX derived from (A) and containing (I); (2) a plasmid vector containing the

XX DNA sequence (III) for (1), under control of a promoter; (3) a helper

XX construct for complementing the coding sequence exchanged by (1); (4)

XX producing (V); (5) producing the plasmids of (2); (6) producing helper

XX constructs of (3); (7) a kit containing the vector of (2) or the helper

XX construct of (3); (8) a DNA molecule (IV) containing at least one coding

XX sequence for (1); (9) a kit containing (IV); (10) a kit for performing

XX methods (5) or (6); (11) a DNA construct that encodes (1) and can persist

XX (12) and is transcribed in target cells, but is preferably not replicable;

XX (13) a recombinant virus (RV), particularly adeno or retro, that encodes

XX (1) and is expressed after infection into a target cell to produce a

XX cytoplasmic replicon that is continuously replenished; (13) producing

XX recombinant DNA viruses or virions having a DNA genome that lacks a

XX specific gene function, in which this function is provided from a

XX recombinant vector system with a RNA genome. The products of the

XX invention have cardioactive activity and can be used for gene therapy.

XX (1) is used to produce gene therapy vectors, particularly plasmids or

XX virions, and these vectors are used for specific transfer to cardiac

XX muscle, for diagnosis, prevention or treatment of cardiac disease, either

XX congenital or acquired. (1) Are also used to complement vectors that lack

XX particular gene sequences, particularly vectors derived from DNA viruses.

XX Vectors based on (1) transfer genes to cardiac myocytes without

XX immunological or other side effects. The RNA genome can replicate,

XX providing efficient gene transfer and long-term expression of the

XX therapeutic gene. CBV is naturally tropic for heart muscle and since it

XX does not produce DNA during its life cycle, overcomes the danger that

CC foreign genes will become integrated in the target cell genome. By using
 CC (II) to replace part of the viral coding region, large (II) sequences may
 CC be accommodated. (I) is easily packaged in CVB capsid proteins.
 XX
 SQ Sequence 35 BP; 14 A; 2 C; 7 G; 12 T; 0 other;

Query Match 14.4%; Score 13; DB 22; Length 35;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 8 aaattattcaaa 20
 |||
 Db 27 AAATATTTCAAA 15

RESULT 16

AAF90560
 ID AAF90560 standard; DNA; 36 BP.

AC AAF90560;

DT 22-AUG-2001 (first entry)

DE Dissociation transposon 3' flanking nested primer 3A.

KW ET1158 gene; GT6839 gene; ET5262 gene; herbicide; screening;

KW herbicide tolerance; transgenic plant; crop protection;

KW Dissociation; transposon; maize; PCR primer; ss.

OS Zea mays.

PN MO200144277-A2.

PD 21-JUN-2001.

PE 14-DEC-2000; 2000MO-EP12748.

PR 16-DEC-1999; 99US-0465040.

PA (SYGN) SYNGENTA PARTICIPATIONS AG.

PI Wegrich Glover L, Budziszewski GJ, Levin JZ, Zhou Q;

DR WPI; 2001-398122/42.

XX New herbicide target genes encoding proteins having ET1158, GT6839 or
 PT ET5262 activity, for identifying an inhibitor of protein activity -
 PS Example 1; Page 39; 67pp; English.

CC The present sequence is that of primer 3A, which is 1 of a set of
 CC 3 nested primers (see also AAF90561 and AAF90562) homologous to
 CC the 3' DS3 region of Dissociation transposon. Arbitrary degenerate
 CC primers (see AAF90551-56) were used to prime Arabidopsis thaliana
 CC genomic DNA flanking the site of a DS transposon insertion. The
 CC degenerate primers were used in combination with 2 sets of 3,
 CC nested, transposon-specific primers homologous to the DS3 region or
 CC DS5 region (see AAF90557-59) of the DS element, which lie at the
 CC outermost ends of the transposon. Low- and high-stringency PCR
 CC amplifications were performed using the TAIL-PCR protocol. DNA
 CC fragments were produced which corresponded to the genomic DNA that
 CC was directly adjacent to the transposon insertion. Sequence
 CC analysis of PCR products from tagged seedling lethal lines ET1158,
 CC GT6839 and ET5262 identified 3 novel genes (see AAF90548-50) each
 CC of which was essential for Arabidopsis seedling growth and
 CC development. The essentiality of the genes provides a means of
 CC discovering new herbicides. Screening assays for identifying
 CC inhibitors that are potential herbicides are provided. The
 CC invention is also applied to the development of herbicide tolerant
 CC plants, and plant tissues, seeds and cells.

XX Sequence 36 BP; 6 A; 4 C; 11 G; 15 T; 0 other;

Query Match 14.4%; Score 13; DB 22; Length 36;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 36 tatgttttcattgt 48
 |||
 Db 21 tatgttttcattgt 33

RESULT 17

AAAT34130/c
 ID AAAT34130 standard; cDNA; 38 BP.

AC AAAT34130;

DT 21-OCT-1996 (first entry)

DE I3L promoter-HTLV-I fusion primer MM116.

KW Avipox virus; poxvirus; vector; vaccine; attenuation; HTLV-I

KW human lymphotropic virus; safety; ALVAC; NYVAC; primer; PCR;

KW polymerase chain reaction; vaccinia virus; retrovirus; ss.

OS Synthetic.

PN MO9621727-A1.

PD 18-JUL-1996.

PE 16-JAN-1996; 96MO-US00547.

PR 13-JAN-1995; 95US-0372664.

PA (VIRO-) VIROGENETICS CORP.

PI Franchini G, Gallo RC, Paolletti E, Tartaglia J;

DR WPI; 1996-342282/34.

XX Attenuated recombinant pox-viruses expressing HTLV antigens - for
 PT safe vaccination against HTLV infection, and for therapy and
 PT diagnosis
 PS Example 12; Page 96; 165pp; English.

CC Primers MM093 (AAAT34127) and MM110 (AAAT34128) were used to generate
 CC a 100 bp PCR fragment, PCR-HTLV18, contg. the I3L promoter fused to
 CC the 5'-end of the HTLV-I envelope gene, using pMM102 as template.
 CC Primers MM113 (AAAT34129) and MM116 (AAAT34130) were used to generate a
 CC 1,500 bp PCR fragment, PCR-HTLV21, contg. the 3' end of the I3L
 CC promoter fused to the entire HTLV-I envelope gene, using p17-SST as
 CC template. MM093 and MM116 were then used as to amplify PCR-HTLV18
 CC and PCR-HTLV21. The I3L-promoted envelope gene was cloned between
 CC canarypox virus C5 flanking arms and also between vaccinia HA
 CC flanking arms, and a NYVAC recombinant expressing HTLV-I env was
 CC generated. This effectively primed protective immune responses
 CC against retrovirus challenge.

XX Sequence 38 BP; 15 A; 6 C; 8 G; 9 T; 0 other;

Query Match 14.4%; Score 13; DB 17; Length 38;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtaattttat 38
 |||
 Db 25 CTTGTAATTTTAT 13

RESULT 18

AA035630/C
 ID AA035630 standard; DNA; 39 BP.
 XX
 AC AA035630;
 XX
 DT 24-FEB-1993 (first entry)
 XX
 DE HIV-2 env 3' fragment primer HIV2B2.
 XX
 KW NYVAC; recombinant; HIV-2; Copenhagen vaccine; vaccinia virus;
 KM virulence factors; deletion loci; recipient loci; env; amplify; PCR;
 KM polymerase chain reaction; gp120; transfection; Vero cells; envelope;
 KM cell surface; ss.
 XX
 OS Synthetic.
 XX
 PN WO9215672-A.
 PD 17-SEP-1992.
 XX
 PF 09-MAR-1992; 92WO-US01906.
 XX
 PR 07-MAR-1991; 91US-0666056.
 PR 11-JUN-1991; 91US-0713967.
 PR 06-MAR-1992; 92US-0847951.
 XX
 PA (VIRO-) VIROGENETICS CORP.
 XX
 PI Cox WI, De Taisne C, Francis J, Gettig RR, Johnson GP;
 PI Limbach KJ, Norton EK, Paoletti E, Perkus ME, Pincus SE;
 PI Riviere M, Tartaglia J, Taylor J;
 XX
 DR WPI, 1992-331718/40.
 XX
 PT Vaccine comprises recombinant, attenuated pox-virus - use for
 PT vaccinating against viral infections such as rabies, hepatitis B,
 PT HIV, HSV, EBV, CMV, mumps etc.
 PS
 PS Disclosure, Page 156; 455pp; English.
 XX
 CC The sequences given in AA035624-32 were used in the construction of
 CC NYVAC recombinants expressing HIV-2 gene products. NYVAC is a
 CC Copenhagen vaccine strain of vaccinia virus which has been modified by
 CC deletion of six non-essential regions of the genome encoding known or
 CC potential virulence factors. The deletion loci were engineered as
 CC recipient loci for the insertion of foreign genes. The HIV-2 env
 CC sequence was isolated in two portions by polymerase chain reaction
 CC (PCR). These fragments were then ligated also by PCR. The HIV-2
 CC gp120 gene was also isolated by PCR. These HIV-2 genes were
 CC transfected into NYVAC which could then be cultured in Vero cells.
 CC The envelope proteins were found to be present on the cell surface of
 CC cells transformed with the recombinant NYVAC. See AA035501-864.
 CC
 SQ Sequence 39 BP; 15 A; 6 C; 7 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 13; Length 39;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtatattttat 38
 |||||||
 DB 25 CCGTAATTTTAT 13

RESULT 19
 AA035353/C
 ID AA035353 standard; DNA; 39 BP.
 XX
 AC AA035353;
 XX
 DT 18-MAY-1993 (first entry)
 XX

DE PCR primer HIV2B2.
 XX
 KW Human immunodeficiency virus; amplification; env; ISSY strain; ss.
 XX
 OS Synthetic.
 XX
 PN WO9222641-A.
 PD 23-DEC-1992.
 XX
 PF 12-JUN-1992; 92WO-US05107.
 XX
 PR 14-JUN-1991; 91US-0715921.
 PR 11-JUN-1992; 92US-0897382.
 XX
 PA (VIRO-) VIROGENETICS CORP.
 XX
 PI Cox WI, Paoletti E, Tartaglia J;
 PI WPI, 1993-018128/02.
 DR
 XX
 PT Modified recombinant virus with inactivated non-essential genetic
 PT functions - comprises e.g. vaccinia or avipox virus, used as HIV
 PT vaccine
 XX
 PS Example 4; Page 52; 159pp; English.
 XX
 CC The 3' portion of the HIV-2 env gene was derived by PCR. In this
 CC reaction a 270 bp fragment was amplified using oligonucleotides
 CC HIV2B1 and HIV2B2 using PISSY-KPN as template. The prod. was
 CC digested with BamHI and XbaI to yield a 150 bp fragment which was
 CC engineered to contain a T5NT sequence known to be recognised as
 CC vaccinia virus early transcription termination signal, following
 CC the termination codon TAA.
 CC See also AA035328-437.
 CC
 SQ Sequence 39 BP; 15 A; 6 C; 7 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 14; Length 39;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtatattttat 38
 |||||||
 DB 25 CCGTAATTTTAT 13

RESULT 20
 AAA99138
 ID AAA99138 standard; DNA; 39 BP.
 XX
 AC AAA99138;
 XX
 DT 19-JAN-2001 (first entry)
 XX
 DE Plasmid pOMPTTC PCR primer SEQ ID NO:15.
 XX
 KW OMP1 protease; cleavage; fusion protein; membrane protease;
 KW natriferic; Escherichia coli; PCR primer; ss.
 XX
 OS Escherichia coli.
 XX
 PN WO200052193-A1.
 PD 08-SEP-2000.
 XX
 PF 03-MAR-2000; 2000WO-JP01309.
 XX
 PR 04-MAR-1999; 99JP-0057731.
 XX
 PA (SUNR) SUNTORY LTD.

XX Okuno K, Yabuta M, Ohsuye K;
 PI WPI: 2000-579291/54.
 DR
 XX Controlled cleavage of peptides by OmpT protease by amino acid
 PT substitution for ensuring cleavage only at desired site in fission of
 PT fusion proteins -
 XX
 PS Example 2; Fig 9; 144pp; Japanese.
 XX
 CC The present invention describes a method for regulating the cleavage
 CC sites of polypeptides by OmpT protease by preventing cleavage at
 CC unwanted sites by converting the amino acid residue at position +1 to
 CC the site to a specifically defined amino acid (where the residue at
 CC position -1 to the site is Lys or Arg), and/or converting the residue
 CC at position -4 and/or -6 to a specifically defined amino acid. Also
 CC described is a method for the fission of a fusion protein to give a
 CC desired polypeptide by cleavage with OmpT protease, where the fusion
 CC protein has a linker peptide inserted between the desired polypeptide
 CC and the other part of the fusion protein. The fusion protein may be
 CC prepared by expression of DNA encoding in a suitable host cell such as
 CC *Escherichia coli*. OmpT protease is a membrane protease of *Escherichia*
 CC *coli* which cleaves peptide chains at a two-residue sites in which both
 CC residues are basic amino acids such as arginine or lysine. The methods
 CC can be used for the efficient preparation of undegraded desired
 CC polypeptides such as natriuretic peptide by OmpT protease cleavage after
 CC recombinant expression as a fusion protein. AAA99127 to AAA99177 and
 CC AAB3946 to AAB24018 represent sequences used in the exemplification of
 CC the present invention.
 CC
 SQ Sequence 39 BP; 8 A; 9 C; 4 G; 18 T; 0 other;

QY Query Match 14.4%; Score 13; DB 21; Length 39;
 Best Local Similarity 100.0%; Pred. NO. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 26 cgtatattttat 38
 15 cgtatattttat 27

RESULT 21
 AAQ48717/C
 ID AAQ48717 standard; DNA; 40 BP.
 XX
 AC AAQ48717;
 XX
 DT 25-MAR-1994 (first entry)
 XX
 DE Insecticidal protein gene 3' primer.
 XX
 KW Caulobacter; plasmid; insecticidal protein; *Bacillus thuringiensis*;
 KW *Bacillus sphaericus*; larva; mosquito; *Culex*; *Anopheles*; *Psorophora*;
 KW *Mansonia*; *Aedes*; ss.
 XX
 OS Synthetic.
 XX
 PN JP05211866-A.
 XX
 PD 24-AUG-1993.
 XX
 PF 05-JUN-1991; 91JP-0160963.
 XX
 PR 06-JUN-1990; 90JP-0148444.
 XX
 PA (SILM-) SILMARAN SO TANABAL.
 XX
 DR WPI: 1993-298916/38.
 XX
 PT Expression of insecticidal protein - by transforming *Caulobacter*
 PT with plasmid contg. gene coding for insecticidal protein

XX Disclosure; Page 24-25; 27pp; Japanese.
 PS
 CC Two primers (AAQ48716-17) are described for the isolation of
 CC *Bacillus thuringiensis israelensis* DNA. The sequence of one
 CC 27-mer primer (AAQ48716), however is missing from the specification.
 CC Caulobacter transformed with a plasmid contg. a gene encoding
 CC insecticidal protein derived from *Bacillus thuringiensis* or
 CC *Bacillus sphaericus* will proliferate in aq. environment.
 CC They may be consumed by larvae of mosquitoes and are lethal to
 CC *Culex*, *Anopheles*, *Psorophora*, *Mansonia* and *Aedes*.
 CC
 SQ Sequence 40 BP; 19 A; 6 C; 6 G; 9 T; 0 other;

QY Query Match 14.4%; Score 13; DB 14; Length 40;
 Best Local Similarity 100.0%; Pred. NO. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 32 ttttatgttttc 44
 15 ttttATGttttTC 3

RESULT 22
 AAZ61322/C
 ID AAZ61322 standard; DNA; 40 BP.
 XX
 AC AAZ61322;
 XX
 DT 19-JUN-2000 (first entry)
 XX
 DE Primer 3A used to construct a synthetic bar gene.
 XX
 KW bar gene; phosphinothricin acetyltransferase; phosphinothricin toxicity;
 KW protein expression; vaccine; haemoglobin; enzyme; primer; ss.
 XX
 OS Synthetic.
 OS Streptomyces hygroscopicus.
 OS
 PN WO200007431-A1.
 XX
 PD 17-FEB-2000.
 XX
 PF 03-AUG-1999; 99WO-US17806.
 XX
 PR 03-AUG-1998; 98US-0095163.
 PR 03-AUG-1998; 98US-0095167.
 PR 15-DEC-1998; 98US-0112257.
 PR 29-APR-1999; 99US-0131611.
 PR 11-JUN-1999; 99US-0138764.
 XX
 PA (RUTP) UNIV RUTGERS STATE NEW JERSEY.
 XX
 PI Maliga P, Kuroda H, Khan MS;
 XX
 DR WPI: 2000-205525/18.
 XX
 PT New recombinant DNA constructs, for expressing high levels of
 PT heterologous protein in plasmids of higher plants, includes promoter, a
 PT leader sequence and a downstream box element -
 XX
 PS Example 7; Page 69; 164pp; English.
 XX
 CC Primers AAZ61218-45 were used to construct a synthetic *Streptomyces*
 CC hygroscopicus bar gene. The bar gene encodes phosphinothricin
 CC acetyltransferase, which provides protection from phosphinothricin
 CC toxicity. The synthetic gene has improved containment and enhanced
 CC expression in plant plasmids, and is used to produce recombinant DNA
 CC constructs of the invention. The specification describes recombinant DNA
 CC constructs for expressing heterologous proteins (e.g. bar gene product)
 CC in the plasmids of higher plants. The DNA constructs comprise a 5'
 CC regulatory region which includes a promoter element, a leader sequence

CC and a downstream box element operably linked to a coding region of the
 CC heterologous protein. The chimeric regulatory region enhances
 CC translational efficiency of an mRNA molecule encoded by the DNA
 CC construct. The DNA constructs are used for producing transformed
 CC monocot and dicot plants having high levels of heterologous protein
 CC expression. They can be used to drive expression of proteins with
 CC agronomic, industrial or pharmaceutical importance, including production
 CC of vaccines, healthcare products like human haemoglobin, industrial or
 CC household enzymes. Plants which can be transformed with the constructs
 CC of the invention include maize, millet, sorghum, sugar cane, rice,
 CC wheat, barley, oat, rye or turf grass.
 CC
 SQ Sequence 40 BP; 19 A; 6 C; 5 G; 10 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 40;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 18 aaagtctactgta 30
 |||
 Db 33 AAAGTTTACTGTA 21

RESULT 23
 AA61323
 ID AA61323 standard; DNA; 40 BP.
 AC AA61323;
 XX
 DT 19-JUN-2000 (first entry)
 XX
 DE Primer 3B used to construct a synthetic bar gene.
 XX
 KW bar gene: phosphinothricin acetyltransferase; phosphinothricin toxicity;
 KW protein expression; vaccine; haemoglobin; enzyme; primer; ss.
 XX
 OS Synthetic.
 OS Streptomyces hygroscopicus.
 XX
 PN WO200007431-A1.
 XX
 PD 17-FEB-2000.
 XX
 PF 03-AUG-1999; 99WO-US17806.
 XX
 PR 03-AUG-1998; 98US-0095163.
 PR 03-AUG-1998; 98US-0095167.
 PR 15-DEC-1998; 98US-0112257.
 PR 29-APR-1999; 99US-0131611.
 PR 11-JUN-1999; 99US-0138764.
 XX
 PA (RUTE) UNIV RUTGERS STATE NEW JERSEY.
 XX
 PI Maliga P, Kuroda H, Khan MS;
 XX
 DR WPI: 2000-205525/18.
 XX
 PT New recombinant DNA constructs, for expressing high levels of
 PT heterologous protein in plastids of higher plants, includes promoter, a
 PT leader sequence and a downstream box element -
 XX
 XX Example 7; Page 69; 164pp; English.
 XX
 CC Primers AA61218-45 were used to construct a synthetic Streptomyces
 CC hygroscopicus bar gene. The bar gene encodes phosphinothricin
 CC acetyltransferase, which provides protection from phosphinothricin
 CC toxicity. The synthetic gene has improved containment and enhanced
 CC expression in plant plastids, and is used to produce recombinant DNA
 CC constructs of the invention. The specification describes recombinant DNA
 CC constructs for expressing heterologous proteins (e.g. bar gene product)
 CC in the plastids of higher plants. The DNA constructs comprise a 5'
 CC regulatory region which includes a promoter element, a leader sequence

CC and a downstream box element operably linked to a coding region of the
 CC heterologous protein. The chimeric regulatory region enhances
 CC translational efficiency of an mRNA molecule encoded by the DNA
 CC construct. The DNA constructs are used for producing transformed
 CC monocot and dicot plants having high levels of heterologous protein
 CC expression. They can be used to drive expression of proteins with
 CC agronomic, industrial or pharmaceutical importance, including production
 CC of vaccines, healthcare products like human haemoglobin, industrial or
 CC household enzymes. Plants which can be transformed with the constructs
 CC of the invention include maize, millet, sorghum, sugar cane, rice,
 CC wheat, barley, oat, rye or turf grass.
 CC
 SQ Sequence 40 BP; 8 A; 5 C; 9 G; 18 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 40;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 18 aaagtctactgta 30
 |||
 Db 28 aaagtctactgta 40

RESULT 24
 AAV51009/C
 ID AAV51009 standard; DNA; 41 BP.
 AC AAV51009;
 XX
 DT 04-JAN-1999 (first entry)
 XX
 DE Maize polymorphic marker S4162/G5-2 DNA.
 XX
 KW Polymorphic marker; allele-specific; primer; probe; amplification;
 KW hybridisation; plant; hybrid certification; genetic contribution;
 KW progeny; back-cross; hybrid; ancestry; maize; ss.
 XX
 OS Zea mays.
 OS
 FH Key Location/Qualifiers
 FT 21
 FT /tag="a"
 FT /replace="g"
 FT /note="polymorphism"
 XX
 PN WO9824796-A1.
 XX
 PD 11-JUN-1998.
 XX
 PF 01-DEC-1997; 97WO-US21782.
 XX
 PR 07-MAR-1997; 97US-0813507.
 PR 02-DEC-1996; 96US-0032069.
 XX
 PA (AFFY-) AFFYMETRIX INC.
 XX
 PI Landry BS, Lemieux B, Murgineux A, Sapolsky RJ;
 XX
 DR WPI: 1998-333252/29.
 XX
 PT Brassica species allele-specific oligonucleotide probes and primers
 PT - useful for plant breeding
 XX
 XX Claim 1; Page 43; 65pp; English.
 XX
 CC This DNA sequence is a region of a Zea mays genome which contains a
 CC polymorphic marker. This sequence can be used in the construction of
 CC allele-specific primers and probes for amplification or hybridisation,
 CC e.g. to determine common or disparate ancestry between 2 or more plants,
 CC to monitor the genetic contribution of an ancestral plant, to trace the
 CC progeny of proprietary plants, in certification of a hybrid plant or to
 CC identify the progeny of a back-crossed plant with an ancestral plant.

XX Sequence 41 BP; 10 A; 9 C; 6 G; 16 T; 0 other;
SQ

Query Match 14.4%; Score 13; DB 19; Length 41;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 aaaaattatttc 17
|||||
DB 36 AAAAAATTATTTC 24

RESULT 25
AAV5101/c
ID AAV51011 standard; DNA; 41 BP.

XX AAV51011;
AC
XX
XX 04-JAN-1999 (first entry)
XX

DE Maize polymorphic marker S4IG2/G4-1 DNA.

XX Polymorphic marker; allele-specific; primer; probe; amplification;
KW hybridisation; plant; hybrid certification; genetic contribution;
XX progeny; back-cross; hybrid; ancestry; maize; ss.

XX Zea mays.

XX Key Location/Qualifiers
FH variation 21
FT /*tag= a
FT /replace= "g"
FT /note= "polymorphism"

XX W09924796-A1.
XX
XX 11-JUN-1998.

XX 01-DEC-1997; 97WO-US21782.
XX
XX
XX 07-MAR-1997; 97US-0813507.
XX
XX 02-DEC-1996; 96US-0032069.

XX (APFY-) AFFYMETRIX INC.
XX
XX
XX Landry BS, Lemieux B, Murigneux A, Sapolsky RJ;
XX
XX WPI; 1998-333252/29.

XX Brassica species allele-specific oligonucleotide probes and primers
XX - useful for plant breeding
XX
XX Claim 1; Page 43; 65pp; English.

XX This DNA sequence is a region of a Zea mays genome which contains a
XX polymorphic marker. This sequence can be used in the construction of
XX allele-specific primers and probes for amplification or hybridisation,
XX e.g. to determine common or disparate ancestry between 2 or more plants,
XX to monitor the genetic contribution of an ancestral plant, to trace the
XX progeny of proprietary plants, in certification of a hybrid plant or to
XX identify the progeny of a back-crossed plant with an ancestral plant.

XX Sequence 41 BP; 12 A; 8 C; 6 G; 15 T; 0 other;
SQ

Query Match 14.4%; Score 13; DB 19; Length 41;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 aaaaattatttc 17
|||||
DB 30 AAAAAATTATTTC 18

RESULT 26
AAT25074/c
ID AAT25074 standard; DNA; 50 BP.

XX AAT25074;
AC
XX
XX 22-OCT-1996 (first entry)
XX

XX Human gene signature HUMGS07214.
DE
XX

XX Gene signature; messenger RNA; mRNA; relative abundance; frequency;
KW human; cloning; mapping; non-biased library; diagnosis; detection;
XX cell typing; abnormal cell function; ss.

OS Homo sapiens.

XX W09514772-A1.
XX
XX 01-JUN-1995.

XX 11-NOV-1994; 94WO-JP01916.
XX
XX 12-NOV-1993; 93JP-0355504.
XX

XX (MATS/) MATSUBARA K.
PA (OKUBO/) OKUBO K.
XX
XX Matsubara K, Okubo K;
XX
XX WPI; 1995-206931/27.

XX Identifying gene signatures in 3'-directed human cDNA library - e.g.
PT for diagnosis of abnormal cell function, by preparing cDNA that
PT reflects relative abundance of corresp. mRNA in specific human
PT tissues

XX Claim 1; Page 1763; 2245pp; Japanese.

XX A single-stranded DNA (or its complementary strand or the corresp.
CC double-stranded DNA) which comprises one of the 7837 "GS" sequences
CC given in AAT19001-T26837 and which is able to hybridise to part of
CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
CC sequences were obtained from 3'-directed cDNA libraries prepared
CC from various human tissues; synthesis of cDNA was initiated from the
CC 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
CC untranslated sequence is unique to a particular mRNA species, almost
CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
CC is constructed so as to reflect accurately the relative abundance of
CC different mRNAs in the particular tissue from which it was derived.
CC The appearance frequency of a given GS in a cDNA library can be
CC determined (esp. using primers and probes derived from the GS
CC sequences) as a means of diagnosing abnormal cell function or for
CC recognising different cell types.

XX Sequence 50 BP; 26 A; 3 C; 3 G; 18 T; 0 other;
SQ

Query Match 14.4%; Score 13; DB 16; Length 50;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 12 tatttcaagttt 24
|||||
DB 43 TATTCCAAGTTT 31

RESULT 27
AAx52184/c
ID AAX52184 standard; DNA; 50 BP.

XX AAX52184;
AC

[illegible]

KW		replication competent double-stranded polynucleotide; ss.
XX		
OS	Synthetic.	
PN	MO9914318-A1.	
XX		
PD	25-MAR-1999.	
XX		
PF	16-SEP-1998;	98WO-US19312.
PR	16-SEP-1997;	97US-0059017.
XX		
PA	(TEKA) UNIV TEXAS SYSTEM.	
XX		
PI	Evans GA;	
DR	WPI: 1999-244029/20.	
XX		
PT	Synthesis of replication competent double-stranded polynucleotides	
PS	Example 4; Fig 5A; 135pp; English.	
XX		
CC	AA52021-212 represent oligonucleotide primers that were used to	
CC	construct a synthetic DNA plasmid sequence synlux4, to demonstrate the	
CC	method of the invention. Within the synlux4 sequence are included the	
CC	sequences of lux A, lux B, the A and B components of the Vidrio fisherit	
CC	luciferase sequence, positions of pUC19 including the origin of	
CC	replication and replication stability sequences, and the promoter and	
CC	coding sequence for tng kanamycin/neomycin phosphotransferase. The	
CC	specification describes a method for the synthesis of replication	
CC	competent double-stranded polynucleotides. The method comprises	
CC	generating a first set of oligonucleotides corresponding to the plus	
CC	strand and a second set corresponding to the minus strand and	
CC	annealing. The method can be used for preparing polynucleotides	
CC	encoding sequences involved in a biochemical pathway. In particular,	
CC	they can be used to produce polynucleotides encoding enzymes,	
CC	e.g. hexokinase, phosphohexose isomerase, phosphofructokinase-1,	
CC	alcoholase, triose-phosphate isomerase, glyceraldehyde-3-phosphate	
CC	dehydrogenase, phosphoglycerate kinase, phosphoglycerate mutase,	
CC	enolase or pyruvate kinase. They can also be used for the preparation	
CC	of viral particles, artificial genomes and artificial genetic systems.	
XX		
SO	Sequence 50 BP; 16 A; 7 C; 8 G; 19 T; 0 other;	
	Query Match	14.4%; Score 13; DB 20; Length 50;
	Best Local Similarity	100.0%; Pred. No. 2.7e+03;
	Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
OY	54 tatgttgagaca 66 Db 33 tatgttgagaca 45	
RESULT 29		
AAAX70046/c		
ID	AAAX70046 standard; RNA: 17 BP.	
AC	AAAX70046;	
DT	28-JUL-1999 (first entry)	
XX		
DE	Human flt1 VEGF receptor hammerhead ribozyme substrate #1341.	
Vascular endothelial growth factor receptor: VEGF receptor: flt-1; flt-1; KDR: hammerhead ribozyme; hairpin ribozyme; cleavage; tumour angiogenesis; psoriasis; Rheumatoid arthritis; ocular disease; fms-like tyrosine kinase 1; kinase insert domain containing receptor; foetal liver kinase 1; ss. Homo sapiens.		
OS	Homo sapiens.	
PN	WO9715662-A2	

XX 01-MAY-1997. 96WO-US17480.
XX 25-OCT-1996; 96WO-US17480.
XX 11-JAN-1996; 96US-0584040.
XX 26-OCT-1995; 95US-0005974.
XX (CHIR) CHIRON CORP.
XX (RIBO-) RIBOZYME PHARM INC.
XX Escobedo J, McSwigen J, Pavco P, Stinchcomb D;
XX WPI; 1997-259017/23.
XX Nucleic acid molecule modulating VEGF receptor(s) gene expression or
XX mRNA stability - useful for treating e.g. tumour angiogenesis,
XX psoriasis, rheumatoid arthritis, etc., in a human patient
XX Claim 4; Page 87; 218pp; English.
XX The present invention describes nucleic acid molecules which modulate
XX the synthesis, expression and/or stability of a mRNA encoding 1 or more
XX receptors of vascular endothelial growth factor (VEGF). A patient
XX (preferably human) having a condition associated with the level of the
XX fms-like tyrosine kinase 1 (flt-1), kinase insert domain containing
XX receptor (KDR) and/or foetal liver kinase 1 (flk-1) (e.g. tumour
XX angiogenesis, ocular diseases, psoriasis and rheumatoid arthritis) can
XX be treated by administering the nucleic acid molecule or the expression
XX vector to the patient. AAX67275 to AAX75752 represent specific examples
XX of nucleic acid molecules from the present invention.
XX Sequence 17 BP; 8 A; 1 C; 4 G; 4 U; 0 other;

Query Match 13.3%; Score 12; DB 18; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 13 attcaaatgtt 24
DB 15 ATTTCAAAGTTT 4

RESULT 30
AAX70047/C
ID AAX70047 standard; RNA; 17 BP.
XX
XX AAX70047;
XX
XX 28-JUL-1999 (first entry)
XX
XX Human flt1 VEGF receptor hammerhead ribozyme substrate #1342.
XX
XX Vascular endothelial growth factor receptor; flt-1;
XX flk-1; KDR; hammerhead ribozyme; cleavage;
XX tumour angiogenesis; psoriasis; rheumatoid arthritis; ocular disease;
XX fms-like tyrosine kinase 1; kinase insert domain containing receptor;
XX foetal liver kinase 1; ss.
XX
XX Homo sapiens.
XX
XX WO9715662-A2.
XX
XX 01-MAY-1997.
XX
XX 25-OCT-1996; 96WO-US17480.
XX
XX 11-JAN-1996; 96US-0584040.
XX 26-OCT-1995; 95US-0005974.
XX
XX (CHIR) CHIRON CORP.
XX (RIBO-) RIBOZYME PHARM INC.

XX Escobedo J, McSwigen J, Pavco P, Stinchcomb D;
XX WPI; 1997-259017/23.
XX Nucleic acid molecule modulating VEGF receptor(s) gene expression or
XX mRNA stability - useful for treating e.g. tumour angiogenesis,
XX psoriasis, rheumatoid arthritis, etc., in a human patient
XX Claim 4; Page 87; 218pp; English.

XX The present invention describes nucleic acid molecules which modulate
XX the synthesis, expression and/or stability of a mRNA encoding 1 or more
XX receptors of vascular endothelial growth factor (VEGF). A patient
XX (preferably human) having a condition associated with the level of the
XX fms-like tyrosine kinase 1 (flt-1), kinase insert domain containing
XX receptor (KDR) and/or foetal liver kinase 1 (flk-1) (e.g. tumour
XX angiogenesis, ocular diseases, psoriasis and rheumatoid arthritis) can
XX be treated by administering the nucleic acid molecule or the expression
XX vector to the patient. AAX67275 to AAX75752 represent specific examples
XX of nucleic acid molecules from the present invention.

Sequence 17 BP; 8 A; 1 C; 3 G; 5 U; 0 other;

Query Match 13.3%; Score 12; DB 18; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 13 attcaaatgtt 24
DB 14 ATTTCAAAGTTT 3

RESULT 31
AAA21453
ID AAA21453 standard; RNA; 17 BP.

XX
XX AAA21453;
XX

DT 19-JUN-2000 (first entry)

DE Integrin alpha 6 subunit substrate sequence SEQ ID NO:4679.

XX Human; aryl hydrocarbon nuclear transport; ARNT; TIE-2; angiogenesis;
XX integrin alpha 6 subunit; integrin subunit beta 3; hairpin ribozyme;
XX hammerhead ribozyme; angiogenic factor; cytoskeletal; antidiabetic;
XX ophthalmologic; antiinflammatory; antiarthritic; antipsoriatic; ARMD;
XX dermatologic; RNA cleavage; cancer; diabetic retinopathy; arthritis;
XX age related macular degeneration; inflammation; neovascular glaucoma;
XX myopic degeneration; psoriasis; verruca vulgaris; angiofibroma;
XX tuberous sclerosis; pot-wine stain; Sturge Weber syndrome;
XX Kippel-Trenaunay-Weber syndrome; Osler-Weber-Rendu syndrome; ss.

XX Homo sapiens.

XX WO950403-A2.

XX 07-OCT-1999.

XX 24-MAR-1999; 99WO-US06507.

XX 27-MAR-1998; 98US-0079678.

XX (RIBO-) RIBOZYME PHARM INC.

XX Pavco PA, Roberts E, Jarvis T, Coesholt C, McSwigen JA;

XX WPI; 1999-591315/50.

XX Novel ribozymes for modulating the synthesis, expression and/or
XX stability of an mRNA encoding an angiogenic factors

PS Claim 55; Page 209; 305pp; English.

XX The present invention describes enzymatic cleave RNA molecules with
CC RNA cleaving activity, which specifically cleave RNA encoded by an aryl
CC hydrocarbon nuclear transporter (ARNT) gene, an integrin subunit beta 3
CC gene, an integrin alpha 6 subunit gene, or a Tie-2 gene. AAA16775 to
CC AAA17167 and AAA17561 to AAA17622 represent ribozyme sequences for ARNT,
CC and AAA17168 to AAA17560 and AAA17623 to AAA17684 represent their
CC corresponding target sequences. AAA17685 to AAA18385 and AAA19087 to
CC AAA19154 represent ribozyme sequences for Tie-2, and AAA18386 to AAA19086
CC and AAA19155 to AAA19222 represent their corresponding target sequences;
CC AAA19223 to AAA20361 and AAA21501 to AAA21595 represent ribozyme
CC sequences for integrin alpha 6 subunit, and AAA20362 to AAA21500 and
CC AAA21596 to AAA21688 represent their corresponding target sequences;
CC AAA21689 to AAA22475 and AAA23263 to AAA23342 represent ribozyme sequence
CC for integrin subunit beta 3, and AAA22476 to AAA23262, AAA23343 to
CC AAA23422 represent their corresponding target sequences. The ribozymes of
CC the invention are used for modulating the synthesis, expression and/or
CC stability of an mRNA encoding angiogenic factor, especially ARNT,
CC integrin subunit beta-3, integrin subunit alpha-6, or Tie-2. They are
CC especially used to treat cancer, diabetic retinopathy, age related
CC macular degeneration (ARMD), inflammation, and arthritis, as well as
CC neovascular glaucoma, myopic degeneration, psoriasis, verruca vulgaris,
CC angiofibroma of tuberous sclerosis, pot-wine stains, Sturge Weber
CC syndrome, Kippel-Trenaunay-Weber syndrome, Osler-Weber-Rendu syndrome,
CC and other syndromes and diseases related to the levels of ARNT, Tie-2,
CC integrin subunit alpha-6, or integrin subunit beta-3.

XX Sequence 17 BP; 6 A; 1 C; 2 G; 8 U; 0 other;

SO

Query Match 13.3%; Score 12; DB 20; Length 17;
Best Local Similarity 58.3%; Pred. No. 8.3e+03;
Matches 7; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

OY 7 aaattatttca 18
|||||:::||||
Db 4 aaauuuuuuca 15

RESULT 32
AAA21454
ID AAA21454 standard; RNA; 17 BP.
XX
AC AAA21454;
XX
DT 19-JUN-2000 (first entry)
DE
DE Integrin alpha 6 subunit substrate sequence SEQ ID NO:4680.
XX
XX Human: aryl hydrocarbon nuclear transport; ARNT; Tie-2; angiogenesis;
KW Integrin alpha 6 subunit; Integrin subunit beta 3; hairpin ribozyme;
KW hammerhead ribozyme; angiogenic factor; cytoskeletal; antidiabetic;
KW ophthalmologic; antiinflammatory; antiarthritic; antipsoriatic; ARMD;
KW dermatologic; RNA cleavage; cancer; diabetic retinopathy; arthritis;
KW age related macular degeneration; inflammation; neovascular glaucoma;
KW myopic degeneration; psoriasis; verruca vulgaris; angiofibroma;
KW tuberous sclerosis; pot-wine stain; Sturge Weber syndrome;
KW Kippel-Trenaunay-Weber syndrome; Osler-Weber-Rendu syndrome; ss.
KM
XX Homo sapiens.
OS
XX W09950403-A2.
PN
XX 07-OCT-1999.
PD
XX 24-MAR-1999; 99WO-US06507.
PE
XX 27-MAR-1998; 98US-0079678.
PR
XX (RIBO-) RIBOZYME PHARM INC.
PA
XX Pavco PA, Roberts E, Jarvis T, Coesholt C, McSwiggen JA;
PI

XX WPI; 1999-591315/50.

DR Novel ribozymes for modulating the synthesis, expression and/or
XX stability of an mRNA encoding an angiogenic factors
PT
PT Claim 55; Page 209; 305pp; English.

PS

XX The present invention describes enzymatic cleave RNA molecules with
CC RNA cleaving activity, which specifically cleave RNA encoded by an aryl
CC hydrocarbon nuclear transporter (ARNT) gene, an integrin subunit beta 3
CC gene, an integrin alpha 6 subunit gene, or a Tie-2 gene. AAA16775 to
CC AAA17167 and AAA17561 to AAA17622 represent ribozyme sequences for ARNT,
CC and AAA17168 to AAA17560 and AAA17623 to AAA17684 represent their
CC corresponding target sequences. AAA17685 to AAA18385 and AAA19087 to
CC AAA19154 represent ribozyme sequences for Tie-2, and AAA18386 to AAA19086
CC and AAA19155 to AAA19222 represent their corresponding target sequences;
CC AAA19223 to AAA20361 and AAA21501 to AAA21595 represent ribozyme
CC sequences for integrin alpha 6 subunit, and AAA20362 to AAA21500 and
CC AAA21596 to AAA21688 represent their corresponding target sequences;
CC AAA21689 to AAA22475 and AAA23263 to AAA23342 represent ribozyme sequence
CC for integrin subunit beta 3, and AAA22476 to AAA23262, AAA23343 to
CC AAA23422 represent their corresponding target sequences. The ribozymes of
CC the invention are used for modulating the synthesis, expression and/or
CC stability of an mRNA encoding angiogenic factor, especially ARNT,
CC integrin subunit beta-3, integrin subunit alpha-6, or Tie-2. They are
CC especially used to treat cancer, diabetic retinopathy, age related
CC macular degeneration (ARMD), inflammation, and arthritis, as well as
CC neovascular glaucoma, myopic degeneration, psoriasis, verruca vulgaris,
CC angiofibroma of tuberous sclerosis, pot-wine stains, Sturge Weber
CC syndrome, Kippel-Trenaunay-Weber syndrome, Osler-Weber-Rendu syndrome,
CC and other syndromes and diseases related to the levels of ARNT, Tie-2,
CC integrin subunit alpha-6, or integrin subunit beta-3.

XX Sequence 17 BP; 7 A; 1 C; 2 G; 7 U; 0 other;

SO

Query Match 13.3%; Score 12; DB 20; Length 17;
Best Local Similarity 58.3%; Pred. No. 8.3e+03;
Matches 7; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

OY 7 aaattatttca 18
|||||:::||||
Db 3 aaauuuuuuca 14

RESULT 33
AAA21455
ID AAA21455 standard; RNA; 17 BP.
XX
AC AAA21455;
XX
DT 19-JUN-2000 (first entry)
DE
DE Integrin alpha 6 subunit substrate sequence SEQ ID NO:4681.
XX
XX Human: aryl hydrocarbon nuclear transport; ARNT; Tie-2; angiogenesis;
KW Integrin alpha 6 subunit; Integrin subunit beta 3; hairpin ribozyme;
KW hammerhead ribozyme; angiogenic factor; cytoskeletal; antidiabetic;
KW ophthalmologic; antiinflammatory; antiarthritic; antipsoriatic; ARMD;
KW dermatologic; RNA cleavage; cancer; diabetic retinopathy; arthritis;
KW age related macular degeneration; inflammation; neovascular glaucoma;
KW myopic degeneration; psoriasis; verruca vulgaris; angiofibroma;
KW tuberous sclerosis; pot-wine stain; Sturge Weber syndrome;
KW Kippel-Trenaunay-Weber syndrome; Osler-Weber-Rendu syndrome; ss.
KM
XX Homo sapiens.
OS
XX W09950403-A2.
PN
XX 07-OCT-1999.
PD
XX 24-MAR-1999; 99WO-US06507.
PE
XX

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XX 27-MAR-1998: 98US-0079678.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX
XX Pavco PA, Roberts E, Jarvis T, Coesholt C, McSwiggen JA;
XX WPI: 1999-591315/50.
XX
XX Novel ribozymes for modulating the synthesis, expression and/or
XX stability of an mRNA encoding an angiogenic factors -
XX
XX Claim 55; Page 209; 305pp; English.
XX
XX The present invention describes enzymatic nucleic acid molecules with
XX RNA cleaving activity, which specifically cleave RNA encoded by an aryl
XX hydrocarbon nuclear transporter (ARNT) gene, an integrin subunit beta 3
XX gene, an integrin alpha 6 subunit gene, or a Tie-2 gene. AAA16775 to
XX AAA17167 and AAA17561 to AAA17622 represent ribozyme sequences for ARNT,
XX and AAA17168 to AAA17560 and AAA17623 to AAA17684 represent their
XX corresponding target sequences; AAA17685 to AAA18385 and AAA19087 to
XX AAA19154 represent ribozyme sequences for Tie-2, and AAA18386 to AAA19086
XX and AAA19155 to AAA19222 represent their corresponding target sequences;
XX AAA19223 to AAA20361 and AAA21501 to AAA21595 represent ribozyme
XX sequences for integrin alpha 6 subunit, and AAA20362 to AAA21500 and
XX AAA21596 to AAA21688 represent their corresponding target sequences;
XX AAA21689 to AAA22475 and AAA23263 to AAA23342 represent ribozyme sequences
XX for integrin subunit beta 3, and AAA22476 to AAA23262, AAA23343 to
XX AAA23422 represent their corresponding target sequences. The ribozymes of
XX the invention are used for modulating the synthesis, expression and/or
XX stability of an mRNA encoding angiogenic factor, especially ARNT,
XX integrin subunit beta-3, integrin subunit alpha-6, or Tie-2. They are
XX especially used to treat cancer, diabetic retinopathy, age related
XX macular degeneration (AMD), inflammation, and arthritis, as well as
XX neovascular glaucoma, myopic degeneration, psoriasis, verruca vulgaris,
XX angiodioma of tuberous sclerosis, pot-wine stains, Sturge Weber
XX syndrome, Kippel-Trenaunay-Weber syndrome, Osler-Weber-Rendu syndrome,
XX and other syndromes and diseases related to the levels of ARNT, Tie-2,
XX integrin subunit alpha-6, or Integrin subunit beta-3.
XX
XX Sequence 17 BP; 7 A; 1 C; 1 G; 8 U; 0 other;
XX
SQ
Query Match 13.3%; Score 12; DB 20; Length 17;
Best Local Similarity 58.3%; Pred. No. 8.3e+03;
Matches 7; Conservative 5; Mismatches 0; Indels 0; Gaps 0;
QY 7 aaaaattattca 18
| | | | | : | : | : | |
Db 1 aaaaauuuuua 12

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RESULT 34
AAAF03088/C
ID AAF03088 standard; DNA; 17 BP.
XX
XX AAF03088;
XX
XX 16-FEB-2001 (first entry)
XX
XX Hammerhead ribozyme substrate #1383.
XX
XX Ribozyme; erythropoietin; granulocyte colony stimulating factor;
XX interferon alpha; ss.
XX
XX Homo sapiens.
XX
XX WO200061729-A2.
XX
XX 19-OCT-2000.
XX
XX 11-APR-2000; 2000WO-US09721.
XX
XX
XX

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PR 12-APR-1999: 99US-0129390.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX
XX Blatt L, Zwick M, Pavco P, McSwiggen J;
XX WPI: 2000-647423/62.
XX
XX Enzymatic and antisense nucleic acid inhibition of repressor genes,
XX useful for producing e.g. granulocyte colony stimulating factor
XX protein, interferon alpha and erythropoietin -
XX
XX Claim 37; Page 87; 164pp; English.
XX
XX The present invention relates to enzymatic and antisense nucleic acid
XX molecules that act as inhibitors of the expression of repressor genes
XX encoding the TR2 Orphan receptor, EAR3/CODP-TF-1, the GATA
XX transcription factor gene, IRF-2 and/or the CAAAT Displacement
XX protein (CDP). Inhibition of the repressors removes prevents
XX inhibition (and consequently increases expression of) genes involved in
XX the production of erythropoietin, granulocyte colony stimulating factor
XX protein and interferon alpha.
XX
XX Sequence 17 BP; 6 A; 1 C; 0 G; 10 T; 0 other;
XX
SQ
Query Match 13.3%; Score 12; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 4 aaaaaattatt 15
| | | | | : | : | : | |
Db 17 AAAAAAATTATT 6

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RESULT 35
AAAF03089/C
ID AAF03089 standard; DNA; 17 BP.
XX
XX AAF03089;
XX
XX 16-FEB-2001 (first entry)
XX
XX Hammerhead ribozyme substrate #1384.
XX
XX Ribozyme; erythropoietin; granulocyte colony stimulating factor;
XX interferon alpha; ss.
XX
XX Homo sapiens.
XX
XX WO200061729-A2.
XX
XX 19-OCT-2000.
XX
XX 11-APR-2000; 2000WO-US09721.
XX
XX 12-APR-1999; 99US-0129390.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX
XX Blatt L, Zwick M, Pavco P, McSwiggen J;
XX WPI: 2000-647423/62.
XX
XX Enzymatic and antisense nucleic acid inhibition of repressor genes,
XX useful for producing e.g. granulocyte colony stimulating factor
XX protein, interferon alpha and erythropoietin -
XX
XX Claim 37; Page 87; 164pp; English.
XX
XX The present invention relates to enzymatic and antisense nucleic acid
XX molecules that act as inhibitors of the expression of repressor genes
XX encoding the TR2 Orphan receptor, EAR3/CODP-TF-1, the GATA

```

transcription factor gene, IRF-2 and/or the CATT Displacement Protein (CDP). Inhibition of the repressors removes prevents inhibition (and consequently increases expression of) genes involved in the production of erythropoietin, granulocyte colony stimulating factor protein and interferon alpha.

Sequence 17 BP; 8 A; 1 C; 0 G; 8 T; 0 other;

Query Match 13.3%; Score 12; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattatt 15
|||||
DB 14 AAAAAATTATT 3

RESULT 36
AAAF03090/C
ID AAF03090 standard; DNA: 17 BP.

AC AAF03090;
XX 16-FEB-2001 (first entry)
XX

DE Hammerhead ribozyme substrate #1385.

KW Ribozyme; erythropoietin; granulocyte colony stimulating factor;
KM Interferon alpha; ss.

OS Homo sapiens.

XX WO200061729-A2.

XX 19-OCT-2000.

XX 11-APR-2000; 2000WC-US09721.

XX 12-APR-1999; 99US-0129390.

PA (RIBO-) RIBOZYME PHARM INC.

PI Blatt L, Zwick M, Pavco P, MCSwigen J;

WPI: 2000-647423/62.

Enzymatic and antisense nucleic acid inhibition of repressor genes, useful for producing e.g. granulocyte colony stimulating factor protein, interferon alpha and erythropoietin -

Claim 37; Page 87; 164pp; English.

The present invention relates to enzymatic and antisense nucleic acid molecules that act as inhibitors of the expression of repressor genes encoding the TR2 Orphan receptor, EAR3/CODP-TR-1, the GATA transcription factor gene, IRF-2 and/or the CATT Displacement Protein (CDP). Inhibition of the repressors removes prevents inhibition (and consequently increases expression of) genes involved in the production of erythropoietin, granulocyte colony stimulating factor protein and interferon alpha.

Sequence 17 BP; 8 A; 1 C; 0 G; 8 T; 0 other;

Query Match 13.3%; Score 12; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattatt 15
|||||
DB 13 AAAAAATTATT 2

RESULT 37
AAAF03091/C
ID AAF03091 standard; DNA: 17 BP.

AC AAF03091;

DT 16-FEB-2001 (first entry)

DE Hammerhead ribozyme substrate #1386.

KW Ribozyme; erythropoietin; granulocyte colony stimulating factor;
KM Interferon alpha; ss.

OS Homo sapiens.

XX WO200061729-A2.

XX 19-OCT-2000.

XX 11-APR-2000; 2000WC-US09721.

XX 12-APR-1999; 99US-0129390.

PA (RIBO-) RIBOZYME PHARM INC.

PI Blatt L, Zwick M, Pavco P, MCSwigen J;

WPI: 2000-647423/62.

Enzymatic and antisense nucleic acid inhibition of repressor genes, useful for producing e.g. granulocyte colony stimulating factor protein, interferon alpha and erythropoietin -

Claim 37; Page 87; 164pp; English.

The present invention relates to enzymatic and antisense nucleic acid molecules that act as inhibitors of the expression of repressor genes encoding the TR2 Orphan receptor, EAR3/CODP-TR-1, the GATA transcription factor gene, IRF-2 and/or the CATT Displacement Protein (CDP). Inhibition of the repressors removes prevents inhibition (and consequently increases expression of) genes involved in the production of erythropoietin, granulocyte colony stimulating factor protein and interferon alpha.

Sequence 17 BP; 9 A; 0 C; 0 G; 8 T; 0 other;

Query Match 13.3%; Score 12; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattatt 15
|||||
DB 12 AAAAAATTATT 1

RESULT 38
AAA08921

ID AAA08921 standard; DNA: 18 BP.

AC AAA08921;

DT 01-AUG-2000 (first entry)

DE Human survivin DNA antisense oligonucleotide, ISIS 23663.

KW Survivin; inhibitor of apoptosis; IAP; caspase inhibitor; caspase-3;
KM cell cycle regulation; cancer; cytostatic; antisense oligonucleotide; ss.

OS Synthetic.

OS Homo sapiens.

```

FH Key Location/Qualifiers
FT modified_base 1.18
FT /tag= a
FT /note= "phosphorothioate backbone"
PN WO200018781-A1.
XX
XX
XX 06-APR-2000.
XX
XX 23-SEP-1999; 99WO-US22076.
XX
XX 29-SEP-1998; 98US-0163162.
PR 05-APR-1999; 99US-0286407.
XX
XX (ISIS-) ISIS PHARM INC.
XX
XX Bennett CF, Ackermann EJ, Swayze EE, Cowser LM;
PI WPI: 2000-293103/25.
DR
XX Antisense molecules targeted to Survivin, useful for inducing apoptosis
XX in cancer cells
XX
XX Claim 3: Page 66; 73pp; English.
XX
XX This is an antisense oligonucleotide targeted to the coding sequence,
XX nucleotide 393, of human survivin DNA (see AAA08903). AAA08910-49 were
XX analyzed for effect on survivin mRNA levels by quantitative real-time
XX PCR. The data obtained were averages from three experiments. This
XX antisense oligonucleotide provided 0% inhibition of survivin mRNA. It
XX was found that ISIS 23667 (AAA08925) provided 70% inhibition and ISIS
XX 23672 (AAA08930) provided 64% inhibition.
XX Survivin, an IAP (inhibitor of apoptosis) caspase inhibitor,
XX has been found to be involved in cell cycle regulation and is expressed
XX in the G2/M phase of the cell cycle in a cell cycle regulated manner and
XX associates with microtubules of the mitotic spindle. Disruption of this
XX interaction results in loss of survivin's anti-apoptotic function and
XX increased caspase-3 activity during mitosis. Caspase-3 is associated
XX with apoptotic cell death. It is therefore believed that survivin may
XX counteract a default induction of apoptosis in the G2/M phase. It is
XX also believed that the over expression of survivin in cancer may
XX overcome this apoptotic check point, allowing undesired survival and
XX division of cancer cells. Antisense oligonucleotides (ASO's) may be used
XX to down regulate endogenous survivin and to increase caspase-3-dependent
XX apoptosis in cells in the G2/M phase.
XX
XX Sequence 18 BP: 1 A; 3 C; 3 G; 11 T; 0 other:
SQ
Query Match 13.3%; Score 12; DB 21; Length 18;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 51 tcttatgttg 62
DB 1 tcttatgttg 12

```

```

PN WO9402613-A.
XX
XX 03-FEB-1994.
PD
XX
XX 15-JUL-1993; 93WO-EP01861.
PF
XX
XX 17-JUL-1992; 92GB-0015233.
PR
XX
XX (PITM ) PITMAN MOORE INC.
XX
XX Osterhaus ADME, Rimmelzwaan GF, Siebelink CHJ;
PI WPI: 1994-048871/06.
DR
XX
XX Synthetic feline immunodeficiency virus (FIV) polypeptide - for
XX use in vaccines used to combat FIV
XX
XX Example 1: Page 21; 41pp; English.
XX
XX The FIV env gene was amplified by PCR using bone marrow
XX derived DNA, obtained from a FIV-infected cat as source
XX of FIV proviral DNA. The primers used are given in AA062805-06
XX and AA062808-09 and were derived from the nucleotide sequence
XX of the petaluma strain of FIV.
XX
XX Sequence 19 BP: 9 A; 3 C; 5 G; 2 T; 0 other:
SQ
Query Match 13.3%; Score 12; DB 15; Length 19;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 50 tcttatgttg 61
DB 13 ttcctatgttg 2

```

```

RESULT 39
AA062806/c
ID AA062806 standard; DNA; 19 BP.
XX
XX AA062806;
XX
XX 21-JUL-1994 (first entry)
XX
XX Env gene 5' primer.
XX
XX FIV, feline immunodeficiency virus; primer: env; cat; petaluma;
XX amplification; polymerase chain reaction; PCR; vaccine; ss.
XX
XX Synthetic.
XX

```

```

RESULT 40
AA269625/c
ID AA269625 standard; DNA; 19 BP.
XX
XX AA269625;
XX
XX 10-SEP-2001 (first entry)
XX
XX Human diallelic marker upstream amplification primer SEQ ID NO:3981.
XX
XX Human genome; diallelic marker; high density disequilibrium map;
XX genomic map; haplotype; phenotype; polymorphic base; genotyping;
XX haplotyping; hybridisation; identification; characterisation;
XX amplification; single nucleotide polymorphism; SNP; PCR primer;
XX diagnosis; ss.
XX
XX Homo sapiens.
XX
XX WO9954500-A2.
XX
XX 28-OCT-1999.
XX
XX 21-APR-1999; 99WO-IB00822.
XX
XX 21-APR-1998; 98US-0082614.
XX
XX 23-NOV-1998; 98US-0109732.
XX
XX (GEST ) GENSET.
XX
XX Cohen D, Blumenfeld M, Chumakov I;
XX
XX WPI: 2000-013267/01.
XX
XX Novel diallelic markers used to construct a high density disequilibrium
XX map of the human genome
XX

```


PS Claim;8; Page 1080; 2745pp; English.

XX AA65654 to AA69578 represent human biallelic markers from the present
 CC invention, which contain a polymorphic base at position 24 of their
 CC nucleotide sequences. AA69579 to AA77440 represent amplification
 CC primers for the biallelic markers. The biallelic markers of the
 CC invention have a variety of uses: they can be used for high density
 CC mapping of the human genome, and in complex association studies and
 CC haplotyping studies which are useful in determining the genetic basis
 CC for disease states. Compositions and methods of the invention can also
 CC be useful for the identification of the targets for the development of
 CC pharmaceutical agents and diagnostic methods, as well as the
 CC characterisation of the differential efficacious responses to and side
 CC effects from pharmaceutical agents acting on a disease as well as other
 CC treatment.

CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
 CC and 3367, are not actually given a sequence in the Sequence Listing
 CC from the present invention.

CC XX Sequence 19 BP; 1 A; 4 C; 5 G; 9 T; 0 other;

SO Query Match 13.3%; Score 12; DB 21; Length 19;
 Best Local Similarity 100.0%; Pred. No. 8.3e+03;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 73 ccaatgaagca 84
 |||||||||
 DB 18 CCAATGAAGCA 7

RESULT 41
 AA75126/C
 ID AA75126 standard; DNA; 19 BP.

XX AA75126;
 DT 10-SEP-2001 (first entry)

XX Human biallelic marker downstream amplification primer SEQ ID NO:9482.

DE Human genome; biallelic marker; high density disequilibrium map;
 KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
 KW haplotyping; hybridisation; identification; characterisation;
 KW amplification; single nucleotide polymorphism; SNP; PCR primer;
 KW diagnosis; ss.

XX Homo sapiens.
 OS
 XX WO954500-A2.
 PN
 XX 28-OCT-1999.
 PD
 XX 21-APR-1999; 99WO-IB00822.
 PF
 XX 21-APR-1998; 98US-0082614.
 PR 23-NOV-1998; 98US-0109732.
 PR
 XX (GEST) GENSET.
 PA
 XX Cohen D, Blumenfeld M, Chumakov I;
 PI
 XX WPI: 2000-013267/01.
 DR
 XX Novel biallelic markers used to construct a high density disequilibrium
 PT map of the human genome -
 XX
 XX Claim 8; Page 2252; 2745pp; English.

XX AA65654 to AA69578 represent human biallelic markers from the present
 CC invention, which contain a polymorphic base at position 24 of their
 CC nucleotide sequences. AA69579 to AA77440 represent amplification
 CC primers for the biallelic markers. The biallelic markers of the

CC invention have a variety of uses: they can be used for high density
 CC mapping of the human genome, and in complex association studies and
 CC haplotyping studies which are useful in determining the genetic basis
 CC for disease states. Compositions and methods of the invention can also
 CC be useful for the identification of the targets for the development of
 CC pharmaceutical agents and diagnostic methods, as well as the
 CC characterisation of the differential efficacious responses to and side
 CC effects from pharmaceutical agents acting on a disease as well as other
 CC treatment.

CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
 CC and 3367, are not actually given a sequence in the Sequence Listing
 CC from the present invention.

CC XX Sequence 19 BP; 7 A; 3 C; 4 G; 5 T; 0 other;

SO Query Match 13.3%; Score 12; DB 21; Length 19;
 Best Local Similarity 100.0%; Pred. No. 8.3e+03;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 21 gttactgtaat 32
 |||||||||
 DB 16 GTTACTGTAAT 5

RESULT 42
 AA005907
 ID AA005907 standard; DNA; 20 BP.

XX AA005907;
 DT 10-JAN-1991 (first entry)

XX HIV mRNA translation inhibiting antisense oligonucleotide (d).
 DE Modified antisense oligonucleotide; HIV mRNA translation inhibition;
 KW HIV-1; transactivator region; leader sequence; tat gene; probe; ss.
 KW Synthetic.

XX EP386563-A.
 PN
 XX 12-SEP-1990.
 PD
 XX 25-FEB-1990; 90EP-0103641.
 PF
 XX 09-MAR-1989; 89DE-3907562.
 PR
 XX (FARB) BAYER AG.
 PA
 XX Stropp U, Baumgarten J, Lobberding A, Springer W, Piel N;
 PI Kretschmer, Kolbl H, Frommer W;
 PI
 XX WPI: 1990-276634/37.
 DR
 XX Chemically modified antisense oligo-nucleotide(s) - inhibiting
 PT translation of HIV mRNA
 PT
 XX Claim 3(d); Page 8; 15pp; German.

XX The chemically modified sequences are opposed to the HIV-1
 CC transactivator region, leader sequences (nt 21-53, 74-161, 202-279)
 CC or exon 2 or 3 of the tat gene (nt 5369-5403, 5421-5548, 5583-5617,
 CC 7967-8366, 8385-9183). This sequence comprises nt 5493-5512
 CC and gives 85% inhibition of HIV-1 tat translation in vitro
 CC at concns. of 5-25 microm.

CC The sequence is useful for treating HIV infections and their
 CC complementary sequences can be used as probes to detect HIV.
 CC Chemical modification comprises replacing one or more internit
 CC phosphodiester linkages by phosphorothioate or methylphosphonate
 CC linkages.
 CC See also AA005904-11.

SQ Sequence 20 BP; 8 A; 4 C; 3 G; 5 T; 0 other;

Query Match 13.3%; Score 12; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 74 caatgaagca 85
|||||
Db 2 caatgaagca 13

RESULT 43

AAT80813/C
ID AAT80813 standard; cDNA; 20 BP.

XX AAT80813;

XX 14-APR-1998 (first entry)

XX Staphylococcus aureus Gene #22 PCR primer sequence 4.

XX Staphylococcus aureus WCUN 29; antagonist; antibacterial; immunogen;

XX vaccine; disease; protection; isolation; PCR primer; ss.

XX Synthetic.

OS Staphylococcus aureus.

XX WO9731114-A2.

XX 28-AUG-1997.

XX 25-FEB-1997; 97WO-GB00524.

XX 26-FEB-1996; 96GB-0004045.

XX (SMRK) SMITHKLINE BEECHAM PLC.

XX Burnham MKR, Hodgson JE;

XX WPI; 1997-435166/40.

XX New Staphylococcus aureus polynucleotide and polypeptide(s) - for
PT isolating antagonist of the polypeptide(s) useful as anti-bacterials
XX

PS Disclosure: Page 52; 117pp; English.

XX The present sequence represents a PCR primer used in the present
CC invention describing novel polypeptides, which can optionally be
CC expressed in NCIMB 40771. The polypeptides, and polynucleotides encoding
CC it, are derived from Staphylococcus aureus. Cells expressing ligands
CC binding the polypeptides can be used to isolated candidate compounds
CC that bind and inhibit the activity of the polypeptides. Such compounds
CC can be used as anti-bacterial compounds. The polypeptides may also be
CC used as immunogens to vaccinate an animal for protection against
CC Staphylococcus aureus caused disease.

XX Sequence 20 BP; 11 A; 4 C; 2 G; 3 T; 0 other;

Query Match 13.3%; Score 12; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcatttc 50
|||||
Db 20 GTTTCATCTTT 9

RESULT 44

AAV72697/C
ID AAV72697 standard; DNA; 20 BP.

XX

AC AAV72697;

XX 17-FEB-1999 (first entry)

XX Corn kernel oil concentration controlling loci marker s1384 primer 2.

XX Corn; kernel oil; concentration; trait controlling loci; genetic marker;

KW Zea mays; breeding; PCR primer; ss.

XX Synthetic.

OS Zea mays.

XX WO9842870-A1.

XX 01-OCT-1998.

XX 19-MAR-1998; 98WO-US05550.

XX 24-MAR-1997; 97US-0041515.

XX (DUPO) DU PONT DE NEMOURS & CO E I.

XX Reltier RS;

XX WPI; 1998-609896/51.

PT Breeding corn with increased oil concentration - comprises use of
PT genetic markers to identify trait loci controlling kernel oil
PT concentration

XX Example 2; Page 5; 50pp; English.

XX A new method has been developed of breeding for corn with increased
CC kernel oil concentration. The method comprises: (a) selecting a corn
CC plant from a breeding population using at least one of the genetic
CC markers s1375, s1384, s1394, s1416, s1422, s1432, s1457, s1480, s1476,
CC s1478, s1484, s1500, s1513, s1529, s1544, s1545, s1630, s1647,
CC s1750, s1756, s1757, s1767, s1772, s1774, s1780, s1797, s1816,
CC s1817, s1836, s1853, s1860, s1870, s1921, s1922, s1925, s1931, s1933,
CC s1939, s1946, s1949, s2054, s2055, s2057, s2058, s2097, s2122, s2125,
CC s2150, s2156, and s2175; and (b) crossing the selected plant with a
CC second plant and obtaining progeny with increased kernel oil
CC concentration. Also described are: (1) a method for identifying corn
CC plants or lines for use as parents to create a breeding population,
CC comprising: (a) genotyping corn plants or lines with one or more of the
CC above genetic markers; and (b) identifying plants or lines which are
CC predicted to produce transgressive segregants for kernel oil
CC concentration; and (2) trait loci controlling kernel oil concentration
CC mapped by the above genetic markers, with the exception of s1480.
CC AAV72694 to AAV72797 represent PCR primers which are used to amplify the
CC genetic markers for use in the method of the invention.

XX Sequence 20 BP; 4 A; 7 C; 3 G; 6 T; 0 other;

Query Match 13.3%; Score 12; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagt 88
|||||
Db 12 TGAAGCAAGTGC 1

RESULT 45

AAV92327
ID AAV92327 standard; DNA; 20 BP.

XX AAV92327;

XX 13-SEP-1999 (first entry)

DE PCR primer used to amplify an ORF of Chlamydia pneumoniae.

XX Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;
 KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;
 KW vaccine; neutralising epitope; PCR primer; ss.

OS Synthetic.
 OS Chlamydia pneumoniae.

PN W09927105-A2.

PD 03-JUN-1999.

PF 20-NOV-1998; 98WO-1B01890.

PR 04-NOV-1998; 98US-0107078.

PR 21-NOV-1997; 97ER-0014673.

XX (GSEST) GENSET.

PI Griffiths R;

DR WPI: 1999-357842/30.

XX Genome sequence of Chlamydia pneumoniae

PS Page 1503; Disclosure; 1912pp; English.

CC AAY91991-X97517 represent PCR primers used to amplify open reading
 CC frames and other nucleic acid sequences from the genome of
 CC Chlamydia pneumoniae (see AAY91990). C. pneumoniae causes respiratory
 CC disease such as pneumonia and bronchitis and is thought to be a
 CC contributing factor in heart disease, sarcoidosis, sinusitis, purulent
 CC otitis media, erythema nodosum or pharyngitis. The polypeptides encoded
 CC by the open reading frames of the C. pneumoniae genome (see AAY34584-
 CC AAY35879) can be used in immunogenic compositions as vaccines. Vectors
 CC containing C. pneumoniae nucleotides sequences can also be used as
 CC immunogenic compositions, especially where the vector directs the
 CC expression of a neutralising epitope of C. pneumoniae.

XX Sequence 20 BP; 3 A; 5 C; 5 G; 7 T; 0 other;

Query Match 13.3%; Score 12; DB 20; Length 20;

Best Local Similarity 100.0%; Pred. No. 8.3e+03;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 atgttttcattgt 48

|||||||||||||

DB 9 atgttttcattgt 20

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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:22:33 : Search time 93.51 Seconds
(without alignments)
791.983 Million cell updates/sec

Title: US-09-531-438-3

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Gapop 60.0 , Gapext 60.0

Searched: 351203 seqs, 11323899 residues

Word size : 0

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Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	4.9	29	3	US-08-816-977-12 Sequence 12, Appl
2	16	4.9	36	1	US-08-629-600-16 Sequence 16, Appl
3	15	4.6	27	1	US-08-120-827-66 Sequence 66, Appl
4	15	4.6	27	1	US-08-478-675-66 Sequence 66, Appl
5	15	4.6	30	5	PCT-US92-10792-3 Sequence 3, Appl1
6	15	4.6	32	1	US-08-256-261-29 Sequence 29, Appl
7	15	4.6	32	3	US-08-852-299-29 Sequence 29, Appl
8	14	4.3	18	2	US-09-205-204-20 Sequence 20, Appl
9	14	4.3	29	3	US-08-816-977-12 Sequence 12, Appl
10	14	4.3	30	2	US-08-629-001A-31 Sequence 31, Appl
11	14	4.3	30	4	US-08-642-274D-110 Sequence 110, App
12	14	4.3	36	1	US-08-629-600-16 Sequence 16, Appl
13	14	4.3	37	2	US-08-403-853-8 Sequence 8, Appl1
14	13	4.0	18	3	US-08-847-844A-113 Sequence 113, App
15	13	4.0	18	4	US-08-686-968C-13 Sequence 13, Appl
16	13	4.0	20	3	US-09-288-461-79 Sequence 79, Appl
17	13	4.0	21	3	US-08-691-045-61 Sequence 61, Appl
18	13	4.0	24	3	US-08-672-215-1 Sequence 1, Appl1
19	13	4.0	28	1	US-08-120-827-64 Sequence 64, Appl
20	13	4.0	28	1	US-08-478-675-64 Sequence 64, Appl
21	13	4.0	30	4	US-08-629-001A-79 Sequence 79, Appl
22	13	4.0	30	4	US-08-642-274D-158 Sequence 158, App
23	13	4.0	31	1	US-08-330-638D-5 Sequence 5, Appl1
24	13	4.0	31	2	US-08-906-746A-5 Sequence 5, Appl1
25	13	4.0	36	1	US-08-247-809A-14 Sequence 14, Appl
26	13	4.0	36	2	US-08-711-728-14 Sequence 14, Appl
27	13	4.0	37	2	US-08-097-554A-45 Sequence 45, Appl

28	13	4.0	37	2	US-08-484-575A-10 Sequence 10, Appl
29	13	4.0	37	3	US-08-477-459-10 Sequence 10, Appl
30	13	4.0	37	3	US-08-480-640A-45 Sequence 45, Appl
31	13	4.0	37	3	US-08-479-869-10 Sequence 10, Appl
32	13	4.0	37	3	US-08-295-802-45 Sequence 45, Appl
33	13	4.0	37	3	US-08-486-414-10 Sequence 10, Appl
34	13	4.0	37	3	US-08-488-237A-45 Sequence 45, Appl
35	13	4.0	37	5	PCT-US94-01826A-10 Sequence 10, Appl
36	13	4.0	37	5	PCT-US94-02252A-10 Sequence 10, Appl
37	13	4.0	38	5	PCT-US96-00547-40 Sequence 40, Appl
38	13	4.0	39	1	US-08-105-483-168 Sequence 168, App
39	13	4.0	39	1	US-08-709-209-168 Sequence 168, App
40	13	4.0	39	1	US-08-303-275-56 Sequence 56, Appl
41	13	4.0	39	1	US-08-458-101-168 Sequence 168, App
42	13	4.0	39	2	US-09-028-361A-19 Sequence 19, Appl
43	13	4.0	40	1	US-08-199-507B-39 Sequence 39, Appl
44	13	4.0	40	1	US-08-441-828-39 Sequence 39, Appl
45	13	4.0	41	3	US-08-930-503A-11 Sequence 11, Appl

ALIGNMENTS

RESULT 1
US-08-816-977-12
; Sequence 12, Application US/08816977
; Patent No. 6080400
; GENERAL INFORMATION:
; APPLICANT: Williams, James A.
; APPLICANT: Byrne, Lisa M.
; TITLE OF INVENTION: Prevention And Treatment Of
; TITLE OF INVENTION: Verotoxin-Induced Disease
; NUMBER OF SEQUENCES: 49
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Medlen & Carroll, LLP
; STREET: 220 Montgomery Street, Suite 2200
; CITY: San Francisco
; STATE: California
; COUNTRY: United States of America
; ZIP: 94104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US-08-816-977
; FILING DATE: 13-MAR-1997
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MacKnight, Kamrin T.
; REGISTRATION NUMBER: 38,230
; REFERENCE/DOCKET NUMBER: OPMD-02450
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 705-8410
; TELEFAX: (415) 397-8338
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 29 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-816-977-12

Query Match 4.9%; Score 16; DB 3; Length 29;
Best Local Similarity 100.0%; Pred. No. 3; Ie+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 189 aaaataatttctta 204
|||||

Db 9 AAAAAATATATTTT 24

RESULT 2

US-08-629-600-16

Sequence 16, Application US/08629600

Patent No. 5783196

GENERAL INFORMATION:

APPLICANT: NORIEGA, Fernando

APPLICANT: LEVINE, Myron M.

TITLE OF INVENTION: GDA MUTANTS OF SHIGELLA

TITLE OF INVENTION: AND VACCINES CONTAINING THE SAME

NUMBER OF SEQUENCES: 18

CORRESPONDENCE ADDRESS:

ADDRESSEE: SUGHRUE, MIOM, ZINN, MACPEAK & SEAS

STREET: 2100 Pennsylvania Avenue, N.W., Suite 800

CITY: Washington, D.C.

STATE: D.C.

COUNTRY: U.S.A.

ZIP: 20037

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/629,600

FILING DATE: 9-APR-1996

CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:

NAME: KIT, Gordon

REGISTRATION NUMBER: 30,764

REFERENCE/DOCKET NUMBER: A-6765

TELEPHONE: (202) 293-7060

TELEFAX: (202) 293-7860

INFORMATION FOR SEQ ID NO: 16:

SEQUENCE CHARACTERISTICS:

LENGTH: 36 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA

HYPOTHETICAL: NO

US-08-629-600-16

Query Match 4.9%; Score 16; DB 1; Length 36;

Best Local Similarity 100.0%; Pred. No. 3e+02;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaaaattattattt 204

Db 14 AAAAAATATATTTT 29

RESULT 3

US-08-120-827-66

Sequence 66, Application US/08120827

Patent No. 5525495

GENERAL INFORMATION:

APPLICANT: KEENE, JACK D.

APPLICANT: KING, PETER H.

APPLICANT: LEVINE, TODD

TITLE OF INVENTION: METHODS AND COMPOSITIONS USEFUL IN THE

TITLE OF INVENTION: RECOGNITION, BINDING AND EXPRESSION OF RIBONUCLEIC ACIDS

NUMBER OF SEQUENCES: 101

CORRESPONDENCE ADDRESS:

ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,

ADDRESS: P.C.

STREET: 1755 Jefferson Davis Highway, Fourth Floor

CITY: Arlington

STATE: Virginia

COUNTRY: U.S.A.

ZIP: 22202

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/120,827

FILING DATE: 15-SEP-1993

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Oblon, No. 5525495man F.

REGISTRATION NUMBER: 24,618

REFERENCE/DOCKET NUMBER: 714-158-0 CIP

TELECOMMUNICATION INFORMATION:

TELEPHONE: (703)413-3000

TELEFAX: (703)413-2220

INFORMATION FOR SEQ ID NO: 66:

SEQUENCE CHARACTERISTICS:

LENGTH: 27 base pairs

TYPE: nucleic acid

STRANDEDNESS: unknown

TOPOLOGY: unknown

MOLECULE TYPE: RNA (genomic)

US-08-120-827-66

Query Match

Best Local Similarity 4.6%; Score 15; DB 1; Length 27;

Matches 5; Conservative 10; Mismatches 0; Indels 0; Gaps 0;

OY 197 tattttattttaa 211

Db 12 UAUUUUUUUUUAAA 26

RESULT 4

US-08-478-675-66

Sequence 66, Application US/08478675

Patent No. 5773246

GENERAL INFORMATION:

APPLICANT: KEENE, JACK D.

APPLICANT: KING, PETER H.

APPLICANT: LEVINE, TODD

TITLE OF INVENTION: METHODS AND COMPOSITIONS USEFUL IN THE

TITLE OF INVENTION: RECOGNITION, BINDING AND EXPRESSION OF RIBONUCLEIC ACIDS

NUMBER OF SEQUENCES: 101

CORRESPONDENCE ADDRESS:

ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,

ADDRESS: P.C.

STREET: 1755 Jefferson Davis Highway, Fourth Floor

CITY: Arlington

STATE: Virginia

COUNTRY: U.S.A.

ZIP: 22202

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/478,675

FILING DATE: 07-JUN-1996

CLASSIFICATION: 536

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/120,827

FILING DATE: 15-SEP-1993

ATTORNEY/AGENT INFORMATION:

NAME: Oblon, No. 5773246man F.

Patent No. 6010897

GENERAL INFORMATION:

APPLICANT: Behnke, Detlef

APPLICANT: Schrott, Bernhard

APPLICANT: Albrecht, Sybille

APPLICANT: G hrs, Karl-Heinz

APPLICANT: Hartmann, Manfred

TITLE OF INVENTION: Expression of signal-peptide-free

TITLE OF INVENTION: staphylokinases
NUMBER OF SEQUENCES: 40
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Neave
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/852,299
FILING DATE: 17-MAY-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/256,261
FILING DATE:
INFORMATION FOR SEQ ID NO: 29:
SEQUENCE CHARACTERISTICS:
LENGTH: 32 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "synthetic DNA from
DESCRIPTION: oligonucleotide synthesis"
US-08-852-299-29

Query Match 4.6%; Score 15; DB 3; Length 32;
Best Local Similarity 100.0%; Pred. No. 7.7e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 154 aaaggaataataaa 168
|||||
DB 30 AAAGCAATATATAA 16

RESULT 8
US-09-205-204-20/c
Sequence 20, Application US/09205204
Patent No. 5958772
GENERAL INFORMATION:
APPLICANT: C. Frank Bennett
APPLICANT: Elizabeth J. Ackermann
TITLE OF INVENTION: ANTISENSE MODULATION OF CELLULAR INHIBITOR OF APOPTOSIS-1 EXPRESS
FILE REFERENCE: RTS-0020
CURRENT APPLICATION NUMBER: US/09/205,204
CURRENT FILING DATE: 1998-12-03
NUMBER OF SEQ ID NOS: 47
SEQ ID NO 20
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Antisense oligonucleotide
US-09-205-204-20

Query Match 4.3%; Score 14; DB 2; Length 18;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 160 aaataataaaa 173
|||||
DB 18 AAATAAATAATAA 5

RESULT 9
US-08-816-977-12/c
Sequence 12, Application US/08816977
Patent No. 6080400
GENERAL INFORMATION:
APPLICANT: Williams, James A.
APPLICANT: Byrne, Lisa M.
APPLICANT: Pugh, Charles S.G.
TITLE OF INVENTION: Prevention And Treatment Of
Verotoxin-Induced Disease
NUMBER OF SEQUENCES: 49
CORRESPONDENCE ADDRESS:
ADDRESSEE: Medlen & Carroll, LLP
STREET: 220 Montgomery Street, Suite 2200
CITY: San Francisco
STATE: California
COUNTRY: United States of America
ZIP: 94104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/816,977
FILING DATE: 13-MAR-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: MacKnight, Kamtin T.
REGISTRATION NUMBER: 38,230
REFERENCE/DOCKET NUMBER: OPMD-02450
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 705-8410
TELEFAX: (415) 397-8338
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 29 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-816-977-12

Query Match 4.3%; Score 14; DB 3; Length 29;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataatatttt 202
|||||
DB 22 AAATAATATTTT 9

RESULT 10
US-08-629-001A-31
Sequence 31, Application US/08629001A
Patent No. 5858661
GENERAL INFORMATION:
APPLICANT: Shiloh, Yosef
TITLE OF INVENTION: ATAXIA-TELANGIECTASIA GENE AND ITS
NUMBER OF SEQUENCES: 139
CORRESPONDENCE ADDRESS:
ADDRESSEE: Kohn & Associates
STREET: 30500 No. 5858661Western Hwy.
CITY: Farmington Hills
STATE: Michigan
COUNTRY: US
ZIP: 48334
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/629,001A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Kohn, Kenneth I.
REGISTRATION NUMBER: 30,955
REFERENCE/DOCKET NUMBER: 2290,00032
TELECOMMUNICATION INFORMATION:
TELEPHONE: (810) 539-5050
TELEFAX: (810) 539-5055
INFORMATION FOR SEQ ID NO: 31:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-629-001A-31

Query Match 4.3%; Score 14; DB 2; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttta 209
|||||
Db 14 ttattttatttta 27

RESULT 11
US-08-642-274D-110
Sequence 110, Application US/08642274D
Patent No. 6200749
GENERAL INFORMATION:
APPLICANT: Shiloh, Yosef
TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
TITLE OF INVENTION: SCREEN FOR A PARTIAL A-T PHENOTYPE
FILE REFERENCE: 229000033
CURRENT APPLICATION NUMBER: US/08/642,274D
CURRENT FILING DATE: 1996-05-03
NUMBER OF SEQ ID NOS: 220
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 110
LENGTH: 30
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: intronic
US-08-642-274D-110

Query Match 4.3%; Score 14; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttta 209
|||||
Db 14 ttattttatttta 27

RESULT 12
US-08-629-600-16/c
Sequence 16, Application US/08629600
Patent No. 5783196
GENERAL INFORMATION:
APPLICANT: NORIEGA, Fernando
APPLICANT: LEVINE, Myron M.
TITLE OF INVENTION: GOA VACCINES OF SHIGELLA
TITLE OF INVENTION: AND VACCINES CONTAINING THE SAME
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:

ADDRESSEE: SUGHRUE, MIOM, ZINN, MACPEAK & SEAS
STREET: 2100 Pennsylvania Avenue, N.W., Suite 800
CITY: Washington, D.C.
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20037

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/629,600
FILING DATE: 9-APR-1996

CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: KIT, Gordon
REGISTRATION NUMBER: 30,764
REFERENCE/DOCKET NUMBER: A-6765
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 293-7060
TELEFAX: (202) 293-7860

INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
HYPOTHETICAL: NO
US-08-629-600-16

Query Match 4.3%; Score 14; DB 1; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttt 202
|||||
Db 27 AAATAATTATTTT 14

RESULT 13
US-08-403-853-8
Sequence 8, Application US/08403853
Patent No. 5844094
GENERAL INFORMATION:
APPLICANT: HUDSON, Peter J.
APPLICANT: KORRT, Alex A.
APPLICANT: IRVING, Robert A.
APPLICANT: ATWELL, John L.
APPLICANT: MALBY, Robyn L.
APPLICANT: POWER, Barbara E.
APPLICANT: COLMAN, Peter M.
TITLE OF INVENTION: TARGET BINDING POLYPEPTIDE
NUMBER OF SEQUENCES: 25
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/403,853
FILING DATE: 30-MAY-1995
CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/AU93/00491
FILING DATE: 24-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: AU PL 4973
FILING DATE: 25-SEP-1992
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16786/189/CHAC
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-403-853-8

Query Match 4.3%; Score 14; DB 2; Length 37;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 83 aaattataattat 96
|||||
Db 1 AATTATTAATTAT 14

RESULT 14

US-08-847-844A-113
Sequence 113, Application US/08847844A
Patent No. 6150160
GENERAL INFORMATION:
APPLICANT: KAZAZIAN JR., HAIG H.
APPLICANT: BOEKE, JEF D.
APPLICANT: MORAN, JOHN V.
APPLICANT: DOMBROSKI, BETH A.
TITLE OF INVENTION: COMPOSITIONS AND METHODS OF USE OF
TITLE OF INVENTION: MAMMALIAN RETROTRANSPOSONS
NUMBER OF SEQUENCES: 137
CORRESPONDENCE ADDRESS:
ADDRESS: PANITCH SCHWARZE JACOBS & NADEL, P.C.
STREET: ONE COMMERCE SQUARE, 2005 MARKET STREET, 22ND FL.
CITY: PHILADELPHIA
STATE: PA
COUNTRY: U.S.A.
ZIP: 19103-7086
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/847,844A
FILING DATE: 28-APR-1997
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/749,805
FILING DATE: 16-NOV-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/006,831
FILING DATE: 16-NOV-1995
ATTORNEY/AGENT INFORMATION:
NAME: DOYLE LEARY Ph.D., KATHRYN
REGISTRATION NUMBER: 36,317
REFERENCE/DOCKET NUMBER: 9596-2302
TELECOMMUNICATION INFORMATION:
TELEPHONE: 215-567-2020
TELEFAX: 215-567-2991

INFORMATION FOR SEQ ID NO: 113:
SEQUENCE CHARACTERISTICS:
LENGTH: 18 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-847-844A-113

Query Match 4.0%; Score 13; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.3e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 150 ttaaaaggaaaa 162
|||||
Db 5 TTAAGAGGAAAA 17

RESULT 15

US-08-686-968C-13
Sequence 13, Application US/08686968C
Patent No. 6221361
GENERAL INFORMATION:
APPLICANT: Cochran, Mark D.
APPLICANT: Junker, David E.
TITLE OF INVENTION: Recombinant Swinepox Virus
FILE REFERENCE: 39119-H/OML
CURRENT APPLICATION NUMBER: US/08/686,968C
CURRENT FILING DATE: 1996-07-25
NUMBER OF SEQ ID NOS: 231
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 13
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer
US-08-686-968C-13

Query Match 4.0%; Score 13; DB 4; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.3e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aattgaattgttaa 243
|||||
Db 5 aattgaattgttaa 17

RESULT 16

US-09-288-461-79
Sequence 79, Application US/09288461
Patent No. 6159694
GENERAL INFORMATION:
APPLICANT: Karas, James G.
TITLE OF INVENTION: Antisense Oligonucleotide Modulation of STAT3
FILE REFERENCE: ISPH-0338
CURRENT APPLICATION NUMBER: US/09/288,461
CURRENT FILING DATE: 1999-04-08
NUMBER OF SEQ ID NOS: 107
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 79
LENGTH: 20
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Sequence
US-09-288-461-79

Query Match 4.0%; Score 13; DB 3; Length 20;

Best Local Similarity 100.0%; Pred. No. 5.2e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 173 attagataaag 185
|||||

Db 2 attagataaag 14

RESULT 17

US-08-691-045-61/c
Sequence 61, Application US/08691045
Patent No. 6015664
GENERAL INFORMATION:
APPLICANT: Henrickson, Kelly J.
TITLE OF INVENTION: VIRUS ASSAY METHOD
NUMBER OF SEQUENCES: 65
CORRESPONDENCE ADDRESS:
ADDRESSEE: Quarles & Brady
STREET: 411 East Wisconsin Avenue
CITY: Milwaukee
STATE: Wisconsin
COUNTRY: U.S.A.
ZIP: 53202-4497
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/691,045
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Baker, Jean C.
REGISTRATION NUMBER: 35,433
REFERENCE/DOCKET NUMBER: 650053,91037
TELECOMMUNICATION INFORMATION:
TELEPHONE: (414) 277-5000
TELEFAX: (414) 271-3552
INFORMATION FOR SEQ ID NO: 61:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: Oligonucleotide
US-08-691-045-61

Query Match 4.0%; Score 13; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 94 tatatgataagta 106
|||||

Db 17 TATATGATAGTA 5

RESULT 18

US-08-672-215-1/c
Sequence 1, Application US/08672215
Patent No. 6020121
GENERAL INFORMATION:
APPLICANT: Ying Bao, Amy Boggs, Pamela R. Contag,
APPLICANT: Nancy A. Federspiel, Alan Herbert,
APPLICANT: Scott J. Hecker, Francois Malouin
TITLE OF INVENTION: INHIBITORS OF REGULATORY PATHWAYS
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700

CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066

COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq for Windows 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/672,215
FILING DATE: June 25, 1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/004,626
FILING DATE: September 29, 1995
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Walburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 219/158
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-672-215-1

Query Match 4.0%; Score 13; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 5.1e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 294 tgtaattttatg 306
|||||

Db 20 TGTATTTTATG 8

RESULT 19

US-08-120-827-64
Sequence 64, Application US/08120827
Patent No. 5525495
GENERAL INFORMATION:
APPLICANT: KEENE, JACK D.
APPLICANT: KING, PETER H.
APPLICANT: LEVINE, TODD
TITLE OF INVENTION: METHODS AND COMPOSITIONS USEFUL IN THE
RECOGNITION, BINDING AND EXPRESSION OF RIBONUCLEIC ACIDS
INVOLVED IN CELL GROWTH, NEOPLASIA AND IMMUNOREGULATION
NUMBER OF SEQUENCES: 101
CORRESPONDENCE ADDRESS:
ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
ADDRESSEE: P. C.
STREET: 1755 Jefferson Davis Highway, Fourth Floor
CITY: Arlington
STATE: Virginia
COUNTRY: U.S.A.
ZIP: 22202
COMPUTER READABLE FORM:
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/120,827
FILING DATE: 15-SEP-1993

```

;      TYPE: nucleic acid
;      STRANDEDNESS: unknown
;

```

```

; TITLE OF INVENTION:  SCREEN
;
; FILE REFERENCE:  2290000333

```

;; CURRENT APPLICATION NUMBER: US/08/642,274D
;; CURRENT FILING DATE: 1996-05-03
;; NUMBER OF SEQ ID NOS: 220
;; SOFTWARE: PatentIn Ver. 2.1
;; SEQ ID NO 158
;; LENGTH: 30
;; TYPE: DNA
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence:intronic
;; OTHER INFORMATION: sequence
US-08-642-274D-158

Query Match 4.0%; Score 13; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 4.9e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 272 aaaaattatttc 284
|||
Db 15 aaaaattatttc 27

RESULT 23
US-08-330-638D-5/C
; Sequence 5, Application US/08330638D
; Patent No. 5731425
; GENERAL INFORMATION:
; APPLICANT: Brizard, Billy
; APPLICANT: Bianca, Darlene
; APPLICANT: Chubert, Richard
; APPLICANT: Vizard, Douglas
; APPLICANT: Hopp, Thomas
; TITLE OF INVENTION: POLYPEPTIDE SURFACE
; TITLE OF INVENTION: MARKER FOR CELLS
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Eastman Kodak Company,
; ADDRESSEE: Patent Legal Staff
; STREET: 343 State Street
; CITY: Rochester
; STATE: New York
; COUNTRY: U.S.A.
; ZIP: 14650-2201
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch,
; MEDIUM TYPE: 1.44 MB storage, (Hewlett Packard)
; COMPUTER: HP Vectra
; OPERATING SYSTEM: MS-DOS Version 6.0
; SOFTWARE: WORD FOR WINDOWS
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/330,638D
; FILING DATE: 28 OCT 1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA: NONE
; ATTORNEY/AGENT INFORMATION:
; NAME: Kierman, Anne B
; REGISTRATION NUMBER: 36,566
; REFERENCE/DOCKET NUMBER: 71255
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (716) 588-2405
; TELEFAX: (716) 477-4646
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31 BASES
; TYPE: NUCLEIC ACID
; STRANDEDNESS: SINGLE
; TOPOLOGY: LINEAR
; MOLECULE TYPE: SYNTHETIC OLIGONUCLEOTIDE
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE: SYNTHETICALLY PREPARED
; IMMEDIATE SOURCE: SYNTHETICALLY PREPARED

;; PUBLICATION INFORMATION: NONE
US-08-330-638D-5

Query Match 4.0%; Score 13; DB 1; Length 31;
Best Local Similarity 100.0%; Pred. No. 4.9e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 124 gtgtctcggggg 136
|||
Db 28 GTGTCTCGGGGG 16

RESULT 24
US-08-906-746A-5/C
; Sequence 5, Application US/08906746A
; Patent No. 5945292
; GENERAL INFORMATION:
; APPLICANT: Brizard, Billy L.
; APPLICANT: Bianca, Darlene W.
; APPLICANT: Chubert, Richard G.
; APPLICANT: Vizard, Douglas L.
; APPLICANT: Hopp, Thomas P.
; TITLE OF INVENTION: Method of Identifying Cells with
; TITLE OF INVENTION: Polypeptide Surface Marker
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Senniger, Powers, Leavitt & Roedel
; STREET: One Metropolitan Square- 16th floor
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/906,746A
; FILING DATE: 06-AUG-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Stone, Paul A.
; REGISTRATION NUMBER: 38,628
; REFERENCE/DOCKET NUMBER: SGM 6874
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 314-231-5400
; TELEFAX: 314-231-4342
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
US-08-906-746A-5

Query Match 4.0%; Score 13; DB 2; Length 31;
Best Local Similarity 100.0%; Pred. No. 4.9e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 124 gtgtctcggggg 136
|||
Db 28 GTGTCTCGGGGG 16

RESULT 25
US-08-247-809A-14/C
; Sequence 14, Application US/08247809A

Patent No. 5569823
GENERAL INFORMATION:
APPLICANT: Peter H. Schreier; Klaus Stenzel; Gunter Adam.
APPLICANT: Edgar Maiss
TITLE OF INVENTION: DEOXYRIBONUCLEIC ACIDS
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: SPRUNG HORN KRAMER & WOODS
STREET: 660 White Plains Road
CITY: Tarrytown
STATE: New York
COUNTRY: U.S.A.
ZIP: 10591-5144
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 2.0 MB
OPERATING SYSTEM: DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/247,809A
FILING DATE: May 23, 1994
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: P 43 178 45.6 (Germany)
FILING DATE: May 28, 1993
ATTORNEY/AGENT INFORMATION:
NAME: Kurt G. Briscoe
REGISTRATION NUMBER: 33,141
REFERENCE/DOCKET NUMBER: Bayer 9049-KGB
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 332-1700
TELEFAX: (914) 332-1844
TELEX:
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-247-809A-14

Query Match 4.0%; Score 13; DB 1; Length 36;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 156 aggaataataa 168
|||||
Db 24 AGGAATAATAA 12

RESULT 26
US-08-711-728-14/c
Sequence 14, Application US/08711728
Patent No. 5973135
GENERAL INFORMATION:
APPLICANT: Peter H. Schreier; Klaus Stenzel; Gunter Adam.
APPLICANT: Edgar Maiss
TITLE OF INVENTION: DEOXYRIBONUCLEIC ACIDS
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: SPRUNG HORN KRAMER & WOODS
STREET: 660 White Plains Road
CITY: Tarrytown
STATE: New York
COUNTRY: U.S.A.
ZIP: 10591-5144
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 2.0 MB
OPERATING SYSTEM: DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/247,809A
FILING DATE: May 23, 1994
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: P 43 178 45.6 (Germany)
FILING DATE: May 28, 1993
ATTORNEY/AGENT INFORMATION:
NAME: Kurt G. Briscoe
REGISTRATION NUMBER: 33,141
REFERENCE/DOCKET NUMBER: Bayer 9049.1-KGB
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 332-1700
TELEFAX: (914) 332-1844
TELEX:
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-711-728-14

OPERATING SYSTEM: DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/711,728
FILING DATE: 03-SEPT-1996
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/247,809
FILING DATE: 23-MAY-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DE 43178456
FILING DATE: 28-MAY-1993
ATTORNEY/AGENT INFORMATION:
NAME: Kurt G. Briscoe
REGISTRATION NUMBER: 33,141
REFERENCE/DOCKET NUMBER: Bayer 9049.1-KGB
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 332-1700
TELEFAX: (914) 332-1844
TELEX:
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-711-728-14

Query Match 4.0%; Score 13; DB 2; Length 36;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 156 aggaataataa 168
|||||
Db 24 AGGAATAATAA 12

RESULT 27
US-08-097-554A-45
Sequence 45, Application US/08097554A
Patent No. 5869312
GENERAL INFORMATION:
APPLICANT: Cochran Ph.D., Mark D
APPLICANT: Junker M.S., David E
TITLE OF INVENTION: Recombinant Swinepox Virus
NUMBER OF SEQUENCES: 112
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/097,554A
FILING DATE: July 22, 1993
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs

```

;
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Swinepox virus
; STRAIN: Kasza
; INDIVIDUAL ISOLATE: S-SPV-001
; IMMEDIATE SOURCE:
; CLONE: 515-85.1
; POSITION IN GENOME:
; MAP POSITION: -23.2
; UNITS: %G
;
US-08-097-554A-45

```

```

Query Match          4.0%; Score 13; DB 2; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 231 aatgaattgttaa 243
    |||||
Db 10 AATGAATTGTAA 22

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RESULT 28
US-08-484-575A-10
; Sequence 10, Application US/08484575A
; Patent No. 5925358
; GENERAL INFORMATION:
; APPLICANT: Mark D. Cochran and David E. Junker
; TITLE OF INVENTION: Recombinant Fowlpox viruses and Uses Thereof
; NUMBER OF SEQUENCES: 42
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: John P. White
; STREET: 1185 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/484,575A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: White Esq, John P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212)278-0450
; TELEFAX: (212)391-0525
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 37 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
;
US-08-484-575A-10

```

```

Query Match          4.0%; Score 13; DB 2; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 231 aatgaattgttaa 243

```

```

Db 10 AATGAATTGTAA 22
    |||||

```

```

RESULT 29
US-08-477-459-10
; Sequence 10, Application US/08477459
; Patent No. 6001369
; GENERAL INFORMATION:
; APPLICANT: Mark D. Cochran
; TITLE OF INVENTION: Recombinant Fowlpox viruses and Uses
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: John P. White
; STREET: 1185 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/477,459
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: White Esq, John P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 278-0400
; TELEFAX: (212) 391-0525
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 37 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
;
US-08-477-459-10

```

```

Query Match          4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 231 aatgaattgttaa 243
    |||||
Db 10 AATGAATTGTAA 22

```

```

RESULT 30
US-08-480-640A-45
; Sequence 45, Application US/08480640A
; Patent No. 6033904
; GENERAL INFORMATION:
; APPLICANT: Cochran, Mark D.
; APPLICANT: Junker, David E.
; TITLE OF INVENTION: Recombinant Swinepox virus
; NUMBER OF SEQUENCES: 225
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: John P. White
; STREET: 1185 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk

```

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA: US/08/480,640A
APPLICATION NUMBER: US/08/480,640A
FILING DATE: 07-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
REGISTRATION NUMBER: 28,678
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 278-0400
TELEFAX: (212) 391-0525
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Swinepox virus
STRAIN: Kasza
INDIVIDUAL ISOLATE: S-SPV-001
IMMEDIATE SOURCE:
CLONE: 515-85.1
POSITION IN GENOME:
MAP POSITION: ~23.2
UNITS: %
US-08-480-640A-45

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 231 aattgaattgtaa 243
DB 10 AATTGAATTGTAA 22

RESULT 31
US-08-479-869-10
Sequence 10, Application US/08479869
Patent No. 6123949
GENERAL INFORMATION:
APPLICANT: Cochran Ph.D., Mark D
TITLE OF INVENTION: Recombinant Fowlpox Virus S-PPV-043 and
TITLE OF INVENTION: Uses Thereof
NUMBER OF SEQUENCES: 20
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/479,869
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/024,156
FILING DATE: 26-FEB-1993
ATTORNEY/AGENT INFORMATION:
NAME: White Esq, John P

TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
US-08-479-869-10

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 231 aattgaattgtaa 243
DB 10 AATTGAATTGTAA 22

RESULT 32
US-08-295-802-45
Sequence 45, Application US/08295802
Patent No. 6127163
GENERAL INFORMATION:
APPLICANT: Cochran Ph.D., Mark D
APPLICANT: Junker M.S., David E
TITLE OF INVENTION: Recombinant Swinepox Virus
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/295,802
FILING DATE: Herewith
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Swinepox virus
STRAIN: Kasza
INDIVIDUAL ISOLATE: S-SPV-001
IMMEDIATE SOURCE:
CLONE: 515-85.1
POSITION IN GENOME:
MAP POSITION: ~23.2

UNITS: %G
US-08-295-802-45

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
Db 10 AATGAATTGTAA 22

RESULT 33
US-08-486-414-10
Sequence 10, Application US/08486414B
Patent No. 6136318
GENERAL INFORMATION:
APPLICANT: Cochran, Mark D.
TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUSES AND USES THEREOF
FILE REFERENCE: 42771D
CURRENT APPLICATION NUMBER: US/08/486,414B
CURRENT FILING DATE: 1995-06-07
NUMBER OF SEQ ID NOS: 46
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 10
LENGTH: 37
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Linker
US-08-486-414-10

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
Db 10 aatgaattgttaa 22

RESULT 34
US-08-488-237A-45
Sequence 45, Application US/08488237A
Patent No. 6251403
GENERAL INFORMATION:
APPLICANT: Cochran, Mark D.
TITLE OF INVENTION: Recombinant Swinepox Virus
NUMBER OF SEQUENCES: 225
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 1185 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/488,237A
FILING DATE: 07-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
REGISTRATION NUMBER: 28,678
TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 278-0400
TELEFAX: (212) 391-0525
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Swinepox virus
STRAIN: Kasza
INDIVIDUAL ISOLATE: S-SPV-001
IMMEDIATE SOURCE:
CLONE: 515-85.1
POSITION IN GENOME:
MAP POSITION: -23.2
UNITS: %G
US-08-488-237A-45

Query Match 4.0%; Score 13; DB 4; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
Db 10 AATGAATTGTAA 22

RESULT 35
PCT-US94-01826A-10
Sequence 10, Application PC/TUS9401826A
GENERAL INFORMATION:
APPLICANT: Syntro Corporation, et al.
TITLE OF INVENTION: Recombinant Fowlpox Virus S-FPV-043 and Uses Thereof
NUMBER OF SEQUENCES: 20
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US94/01826A
FILING DATE: 28-FEB-1994
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: White Esq, John P
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
PCT-US94-01826A-10

Query Match 4.0%; Score 13; DB 5; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
|||||
Db 10 AATGAATTGTAA 22

RESULT 36
PCT-US94-02252A-10

; Sequence 10, Application PC/TUS9402252A
; GENERAL INFORMATION:
; APPLICANT: Syntro Corporation, et al.
; TITLE OF INVENTION: Recombinant Fowlpox Viruses and Uses Thereof
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John P. White
; STREET: 30 Rockefeller Plaza
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10112
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/02252A
; FILING DATE: 28-FEB-1994
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: White Esq, John P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212)977-9550
; TELEFAX: (212)664-0525
; TELEX: 422523
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 37 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; PCT-US94-02252A-10

Query Match 4.0%; Score 13; DB 5; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
|||||
Db 10 AATGAATTGTAA 22

RESULT 37
PCT-US96-00547-40/C

; Sequence 40, Application PC/TUS9600547
; GENERAL INFORMATION:
; APPLICANT: Virogenetics Corporation
; TITLE OF INVENTION: RECOMBINANT POXVIRUS-HTLV, COMPOSITIONS
; NUMBER OF SEQUENCES: 56
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Curtis, Morris & Safford, P.C.
; STREET: 530 Fifth Avenue, 25th Floor
; CITY: New York
; STATE: New York
; COUNTRY: U.S.A.

ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US96/00547
FILING DATE: 12-JAN-1996
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/372,664
FILING DATE: 13-JAN-1995
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2621
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 38 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
PCT-US96-00547-40

Query Match 4.0%; Score 13; DB 5; Length 38;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 293 ctgtaattttat 305
|||||
Db 25 CTGTAATTTTAT 13

RESULT 38
US-08-105-483-168/C
; Sequence 168, Application US/08105483
; Patent No. 5494807
; GENERAL INFORMATION:
; APPLICANT: Paolelli, Enzo
; TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
; NUMBER OF SEQUENCES: 462
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Curtis, Morris & Safford
; ADDRESSEE: C/o William S. Frommer
; STREET: 530 Fifth Avenue
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/105,483
; FILING DATE: 12-AUG-1993
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/847,951
; FILING DATE: 06-MAR-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Frommer, William S.
; REGISTRATION NUMBER: 25,506
; REFERENCE/DOCKET NUMBER: 454310-2400
; TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 840-3333
 TELEFAX: (212) 840-0712
 INFORMATION FOR SEQ ID NO: 168:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 39 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 US-08-105-483-168

Query Match 4.0%; Score 13; DB 1; Length 39;
 Best Local Similarity 100.0%; Pred. No. 4.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 293 ctgtaattttat 305
 |||||||
 Db 25 CTGTAATTTTAT 13

RESULT 39
 US-08-709-209-168/c
 ; Sequence 168, Application US/08709209
 ; Patent No. 5762938
 ; GENERAL INFORMATION:
 ; APPLICANT: Paoletti, Enzo
 ; TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
 ; TITLE OF INVENTION: STRAIN
 ; NUMBER OF SEQUENCES: 462
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Curtis, Morris & Safford
 ; ADDRESSEE: c/o William S. Frommer
 ; STREET: 530 Fifth Avenue
 ; CITY: New York
 ; STATE: NY
 ; COUNTRY: USA
 ; ZIP: 10036
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patentin Release #1.0, Version #1.25
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/709,209
 ; FILING DATE: 21-AUG-1996
 ; CLASSIFICATION: 424
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: US 08/105,483
 ; FILING DATE: 12-AUG-1993
 ; APPLICATION NUMBER: US 07/847,951
 ; FILING DATE: 06-MAR-1992
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Frommer, William S.
 ; REGISTRATION NUMBER: 25,506
 ; REFERENCE/DOCKET NUMBER: 454310-2400
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (212) 840-3333
 ; TELEFAX: (212) 840-0712
 ; INFORMATION FOR SEQ ID NO: 168:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 39 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; US-08-709-209-168

Query Match 4.0%; Score 13; DB 1; Length 39;
 Best Local Similarity 100.0%; Pred. No. 4.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 293 ctgtaattttat 305
 |||||||

Db 25 CTGTAATTTTAT 13

RESULT 40
 US-08-303-275-56/c
 ; Sequence 56, Application US/08303275
 ; Patent No. 5766598
 ; GENERAL INFORMATION:
 ; APPLICANT: Paoletti, Enzo
 ; APPLICANT: Tartaglia, James
 ; APPLICANT: Cox, William I.
 ; TITLE OF INVENTION: IMMUNODEFICIENCY VIRUS RECOMBINANT
 ; TITLE OF INVENTION: POXVIRUS VACCINE
 ; NUMBER OF SEQUENCES: 205
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Curtis, Morris & Safford
 ; ADDRESSEE: c/o William S. Frommer
 ; STREET: 530 Fifth Avenue
 ; CITY: New York
 ; STATE: New York
 ; COUNTRY: USA
 ; ZIP: 10036
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patentin Release #1.0, Version #1.25
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/303,275
 ; FILING DATE:
 ; CLASSIFICATION: 424
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: US 07/897,382
 ; FILING DATE: 11-JUN-1992
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Frommer, William S.
 ; REGISTRATION NUMBER: 25,506
 ; REFERENCE/DOCKET NUMBER: 454310-2420
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (212) 840-3333
 ; TELEFAX: (212) 840-0712
 ; INFORMATION FOR SEQ ID NO: 56:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 39 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; US-08-303-275-56

Query Match 4.0%; Score 13; DB 1; Length 39;
 Best Local Similarity 100.0%; Pred. No. 4.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 293 ctgtaattttat 305
 |||||||
 Db 25 CTGTAATTTTAT 13

RESULT 41
 US-08-458-101-168/c
 ; Sequence 168, Application US/08458101
 ; Patent No. 5766599
 ; GENERAL INFORMATION:
 ; APPLICANT: Paoletti, Enzo
 ; APPLICANT: Perkus, Marion E.
 ; APPLICANT: Taylor, Jill
 ; APPLICANT: Tartaglia, James
 ; APPLICANT: No. 5766599ton, Elizabeth K.
 ; APPLICANT: Riviere, Michel
 ; APPLICANT: de Taisne, Charles
 ; APPLICANT: Limbach, Keith J.
 ; APPLICANT: Johnson, Gerard P.

APPLICANT: Pincus, Steven E.
APPLICANT: Cox, William I.
APPLICANT: Audonnet, Jean-Christophe Francis
APPLICANT: Gettig, Russell Robert
TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
TITLE OF INVENTION: STRAIN
NUMBER OF SEQUENCES: 467
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESSEE: C/O William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/458,101
FILING DATE: 01-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2740
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-458-101-168

Query Match 4.0%; Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 293 ctgtaattttat 305
|||||
DB 25 CTGTAATTTTAT 13

RESULT 42
US-09-028-361A-19
Sequence 19, Application US/09028361A
Patent No. 5962296
GENERAL INFORMATION:
APPLICANT: ETTWILLER, LAURENCE
APPLICANT: XU, SHUANG-YONG
TITLE OF INVENTION: METHOD FOR CLONING AND
TITLE OF INVENTION: PRODUCING THERMOMICROBIUM ROSEUM DNA
TITLE OF INVENTION: POLYMERASE I IN E. COLI
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: New England Biolabs, Inc.
STREET: 32 Tozer Road
CITY: Beverly
STATE: MA
COUNTRY: USA
ZIP: 01915
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq Version 2.0
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/028,361A
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Williams, Gregory D.
REGISTRATION NUMBER: 30901
REFERENCE/DOCKET NUMBER: NEB-134
TELECOMMUNICATION INFORMATION:
TELEPHONE: 978-927-5054
TELEFAX: 978-927-1705
TELEX:
INFORMATION FOR SEQ ID NO: 19:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-09-028-361A-19

Query Match 4.0%; Score 13; DB 2; Length 39;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 62 atataataattat 74
|||||
DB 12 ATATATTAATAT 24

RESULT 43
US-08-199-507B-39/C
Sequence 39, Application US/08199507B
Patent No. 5472841
GENERAL INFORMATION:
APPLICANT: JAYASENA, S. AND GOLD, L.
TITLE OF INVENTION: NUCLEIC ACID LIGAND INHIBITORS
NUMBER OF SEQUENCES: 59
CORRESPONDENCE ADDRESS:
ADDRESSEE: Swanson & Bratschun, L.L.C.
STREET: 8400 E. Prentice Avenue, Suite 200
CITY: Englewood
STATE: Colorado
COUNTRY: USA
ZIP: 80111
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 Inch, 360 KB storage
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/199,507B
FILING DATE: 22 FEBRUARY 1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/714,131
FILING DATE: 10-JUNE-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/536,428
FILING DATE: 11-JUNE-1990
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: NEX13
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 39:
SEQUENCE CHARACTERISTICS:

LENGTH: 40 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-199-507B-39

Query Match 4.0%; Score 13; DB 1; Length 40;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 157 ggaataataaat 169
|||||
DB 22 GGAATAATAAAT 10

RESULT 44
US-08-441-828-39/C
Sequence 39, Application US/08441828
Patent No. 5734034
GENERAL INFORMATION:
APPLICANT: JAYASENA, S. AND GOLD, L.
TITLE OF INVENTION: NUCLEIC ACID LIGAND INHIBITORS
TITLE OF INVENTION: OF HUMAN NEUTROPHIL ELASTASE
NUMBER OF SEQUENCES: 59
CORRESPONDENCE ADDRESS:
ADDRESSEE: Swanson & Bratschun, L.L.C.
STREET: 8400 E. Prentice Avenue, Suite 200
CITY: Englewood
STATE: Colorado
COUNTRY: USA
ZIP: 80111
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44 MB
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/441,828
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/199,507
FILING DATE: 22 FEBRUARY 1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/714,131
FILING DATE: 10-JUNE-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/536,428
FILING DATE: 11-JUNE-1990
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: NEX13
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 39:
SEQUENCE CHARACTERISTICS:
LENGTH: 40 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-441-828-39

Query Match 4.0%; Score 13; DB 1; Length 40;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 157 ggaataataaat 169
|||||

DB 22 GGAATAATAAAT 10

RESULT 45
US-08-930-503A-11
Sequence 11, Application US/08930503A
Patent No. 6132731
GENERAL INFORMATION:
APPLICANT:
TITLE OF INVENTION: MORINE LEUKEMIA VIRUS VECTORS (AS AMENDED)
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: WENDEROTH, LIND & PONACK, L.L.P.
STREET: 2033 K Street, N.W. - Suite 800
CITY: Washington
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20006
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/930,503A
FILING DATE: October 8, 1997
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9506782.3
FILING DATE: 01-APR-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/GB96/00776
FILING DATE: 01-APR-1996
ATTORNEY/AGENT INFORMATION:
NAME: Warren M Cheek, Jr.
REGISTRATION NUMBER: 33,367
REFERENCE/DOCKET NUMBER:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-721-8200
TELEFAX: 202-721-8250
TELEX:
INFORMATION FOR SEQ ID NO: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 41 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-930-503A-11

Query Match 4.0%; Score 13; DB 3; Length 41;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 251 ttccgggggggaat 263
|||||
DB 2 TTCAGGGGGGAAT 14

Search completed: January 24, 2002, 03:24:24
Job time: 3711 sec

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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:19:03 ; Search time 1494.92 Seconds

(without alignments)
3608.606 Million cell updates/sec

Title: US-09-531-438-3

Perfect score: 327
Sequence: 1 atttgggatatcttaattt.....tttcattttttttattgtt 327

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 1472140 seqs, 8248589755 residues

Word size: 0

Total number of hits satisfying chosen parameters: 541028

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

GenEmbl:*

- 1: gb_da:*
- 2: gb_bg:*
- 3: gb_in:*
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- 5: gb_ov:*
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- 8: gb_pl:*
- 9: gb_pr:*
- 10: gb_ro:*
- 11: gb_sts:*
- 12: gb_sy:*
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- 17: em_hum:*
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- 25: em_ro:*
- 26: em_sts:*
- 27: em_sy:*
- 28: em_un:*
- 29: em_vl:*
- 30: em_htgo_hum:*
- 31: em_htgo_inv:*
- 32: em_htgo_rod:*
- 33: em_htg_hum:*
- 34: em_htg_inv:*
- 35: em_htg_rod:*
- 36: em_htg_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	4.9	29	6	AR099868 Sequence
2	16	4.9	32	6	E27913 Method for
3	16	4.9	36	3	CEANONMYR X97533 C.elegans D
4	16	4.9	36	6	AR019036 Sequence
5	15	4.6	27	6	AR014030 Sequence
6	15	4.6	27	6	121980 Sequence 66
7	15	4.6	32	6	AR037189 Sequence
8	14	4.3	29	6	AR076353 Sequence
9	14	4.3	29	6	AR099868 Sequence
10	14	4.3	30	6	AR028182 Sequence
11	14	4.3	30	6	AR138585 Sequence
12	14	4.3	32	6	E27913 Method for
13	14	4.3	36	6	AR019036 Sequence
14	14	4.3	37	6	AR063204 Sequence
15	14	4.3	45	6	AX049973 Sequence
16	14	4.3	45	6	AX049974 Sequence
17	14	4.3	45	6	AX099855 Sequence
18	14	4.3	45	6	AX099856 Sequence
19	14	4.3	45	6	AX137975 Sequence
20	14	4.3	45	6	AX137976 Sequence
21	14	4.3	50	6	AX159856 Sequence
22	13	4.0	20	6	AR146953 Sequence
23	13	4.0	20	6	AR121058 Sequence
24	13	4.0	20	6	AX076045 Sequence
25	13	4.0	23	6	A97479 Sequence 35
26	13	4.0	24	6	AX093544 Sequence
27	13	4.0	24	6	AX164353 Sequence
28	13	4.0	24	12	AB069100 Synthetic
29	13	4.0	25	6	AX042574 Sequence
30	13	4.0	25	6	AX043268 Sequence
31	13	4.0	26	6	AX039624 Sequence
32	13	4.0	26	6	AX039624 Sequence
33	13	4.0	28	6	AR014028 Sequence
34	13	4.0	28	6	121978 Sequence 64
35	13	4.0	29	6	AX012366 Sequence
36	13	4.0	29	6	E59972 Highly acti
37	13	4.0	30	6	AR028230 Sequence
38	13	4.0	30	6	AR138633 Sequence
39	13	4.0	30	6	AX063379 Sequence
40	13	4.0	31	6	195122 Sequence 5
41	13	4.0	33	5	XELARSE59 K01606 Xenopus lae
42	13	4.0	36	6	AA1027 Sequence 14
43	13	4.0	36	6	AR082586 Sequence
44	13	4.0	36	6	AX167671 Sequence
45	13	4.0	36	6	128261 Sequence 14

ALIGNMENTS

RESULT 1
AR099868
LOCUS AR099868 29 bp DNA
DEFINITION Sequence 12 from patent US 6080400.
ACCESSION AR099868
VERSION AR099868.1 GI:12810316
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 29)
AUTHORS Williams,J.A. and Byrne,L.Marie.
TITLE Compositions for the prevention and treatment of verotoxin-induced disease
JOURNAL Patent: US 6080400-A 12 27-JUN-2000;
FEATURES location/Qualifiers
source 1..29
BASE COUNT 11 a 2 c 5 g 11 t
ORIGIN

Query Match 4.9%; Score 16; DB 6; Length 29;
 Best Local Similarity 100.0%; Pred. No. 9.2e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
 |||||||
 DB 9 AAAATAATTATTTT 24

RESULT 2

E27913 32 bp DNA PAT 07-FEB-2001
 LOCUS E27913
 DEFINITION Method for detecting foreign DNA fragment insert in Vero toxin gene.

ACCESSION E27913
 VERSION E27913.1 GI:13020766
 KEYWORDS JP 1999243996-A/3.
 SOURCE unidentified.
 ORGANISM unidentified.

REFERENCE 1 (bases 1 to 32)
 AUTHORS Masahiro, K., Y. N. N. and Kawamura, K. S.
 TITLE Method for detecting foreign DNA fragment insert in Vero toxin gene
 JOURNAL Patent: JP 1999243996-A 3 14-SEP-1999;

COMMENT

OS Unidentified
 PN JP 1999243996-A/3
 PD 14-SEP-1999
 PE 27-FEB-1998 JP 1998047677
 PR

PI MASAHIRO KUSUMOTO, YOSHIAKI NISHIYA, YOSHIOHISA KANAMURA, PI
 KUNIHITO SHINAGAWA
 PC C12Q1/68,C12N15/09/(C12N15/09,C12R1:185),C12N15/00,
 (C12N15/00, PC C12R1:185)
 CC Strandedness: Both;
 CC Topology: Linear;
 FH key
 FT source
 FT Location/Qualifiers
 1.32
 Location/Qualifiers
 /organism="unidentified"
 /db_xref="taxon:32644"

BASE COUNT 12 a 2 c 4 g 14 t
 ORIGIN

Query Match 4.9%; Score 16; DB 6; Length 32;
 Best Local Similarity 100.0%; Pred. No. 9.2e+03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
 |||||||
 DB 4 AAAATAATTATTTT 19

RESULT 3

CEANONYFR 36 bp DNA INV 06-MAY-1997
 LOCUS CEANONYFR
 DEFINITION C. elegans DNA fragment with rearrangement site.

ACCESSION X97532
 VERSION X97532.1 GI:1296496
 KEYWORDS

SOURCE Caenorhabditis elegans.
 ORGANISM Caenorhabditis elegans

Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;
 Rhabditoidae; Rhabditidae; Pelodierinae; Caenorhabditis.

REFERENCE 1 (bases 1 to 36)

AUTHORS Wicky, C., Villeneuve, A. M., Lauper, N., Codourey, L., Tobler, H. and
 Muller, F.

Telomeric repeats (TTAGGC)n are sufficient for chromosome capping

JOURNAL function in Caenorhabditis elegans
 Proc. Natl. Acad. Sci. U.S.A. 93 (17), 8983-8988 (1996)

MEDLINE 96392352

REFERENCE 2 (bases 1 to 36)

AUTHORS Wicky, C.

JOURNAL Direct Submission
 Submitted (24-APR-1996) C. Wicky, University of British Columbia,
 Medical Genetics, 6174 University Boulevard, Vancouver, B.C. V6T
 1Z3, CANADA

FEATURES Location/Qualifiers

source

1.36
 /organism="Caenorhabditis elegans"
 /db_xref="taxon:6239"
 /clone="Bp1"

misc_feature 1.36
 BASE COUNT 9 a 5 c 4 g 18 t
 ORIGIN

Query Match 4.9%; Score 16; DB 3; Length 36;
 Best Local Similarity 100.0%; Pred. No. 9.1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 169 taataattagataaa 184
 |||||||
 DB 23 TAAATTTAGATRAAA 8

RESULT 4

ARO19036 36 bp DNA PAT 05-DEC-1998
 LOCUS ARO19036
 DEFINITION Sequence 16 from patent US 5783196.

ACCESSION ARO19036
 VERSION ARO19036.1 GI:3974150
 KEYWORDS

SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 36)

AUTHORS Noriega, F. R. and Levine, M. M.
 TITLE Gua mutants of shigella spp. and vaccines containing the same
 JOURNAL Patent: US 5783196-A 16 21-JUL-1998;

FEATURES Location/Qualifiers

source 1.36
 /organism="unknown"
 BASE COUNT 11 a 3 c 10 g 12 t
 ORIGIN

Query Match 4.9%; Score 16; DB 6; Length 36;
 Best Local Similarity 100.0%; Pred. No. 9.1e+03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
 |||||||
 DB 14 AAAATAATTATTTT 29

RESULT 5

ARO14030 27 bp DNA PAT 05-DEC-1998
 LOCUS ARO14030
 DEFINITION Sequence 66 from patent US 5773246.

ACCESSION ARO14030
 VERSION ARO14030.1 GI:3971484
 KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 27)

AUTHORS Keene, J. D., Levine, T. and Gao, F.

Methods and compositions useful in the recognition, binding and
 expression of ribonucleic acids involved in cell growth, neoplasia

JOURNAL and immunoregulation
Patent: US 5773246-A 66 30-JUN-1998;
FEATURES Location/Qualifiers
SOURCE 1..27
BASE COUNT 7 a 2 c 2 g 16 t
ORIGIN

Query Match 4.6%; Score 15; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.4e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 197 tattttattttaaa 211
Db 12 TATTTTATTTTAA 26

RESULT 6
LOCUS 121980 27 bp DNA PAT 07-OCT-1996
DEFINITION Sequence 66 from patent US 5525495.
ACCESSION 121980
VERSION 121980.1 GI:1602334
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 27)
AUTHORS Keene,J.D., Levine,T. and Gao,F.
TITLE Methods and compositions useful in the recognition, binding and expression of ribonucleic acids involved in cell growth, neoplasia and immunoregulation
JOURNAL Patent: US 5525495-A 66 11-JUN-1996;
FEATURES Location/Qualifiers
SOURCE 1..27
BASE COUNT 7 a 2 c 2 g 16 t
ORIGIN

Query Match 4.6%; Score 15; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.4e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 197 tattttattttaaa 211
Db 12 TATTTTATTTTAA 26

RESULT 7
LOCUS AR037189/c 32 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 29 from patent US 5801037.
ACCESSION AR037189
VERSION AR037189.1 GI:5955045
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 32)
AUTHORS Behnke,D., Schloft,B., Albrecht,S., Guhrs,K. and Hartmann,M.
TITLE Expression of signal peptide-free scapapylolkinases
JOURNAL Patent: US 5801037-A 29 01-SEP-1998;
FEATURES Location/Qualifiers
SOURCE 1..32
BASE COUNT 3 a 8 c 4 g 17 t
ORIGIN

Query Match 4.6%; Score 15; DB 6; Length 32;
Best Local Similarity 100.0%; Pred. No. 2.4e+04;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 154 aaaggaataataa 168
Db 30 AAAGGAATAATAA 16

RESULT 8
LOCUS AR076353/c 18 bp DNA PAT 30-AUG-2000
DEFINITION Sequence 20 from patent US 5958772.
ACCESSION AR076353
VERSION AR076353.1 GI:10003099
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 18)
AUTHORS Bennett,C.Frank, Ackermann,E.J. and Cowser,L.M.
TITLE Antisense inhibition of cellular inhibitor of apoptosis-1 expression
JOURNAL Patent: US 5958772-A 20 28-SEP-1999;
FEATURES Location/Qualifiers
SOURCE 1..18
BASE COUNT 3 a 2 c 1 g 12 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 6.2e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 160 aaataataataaa 173
Db 18 AAATAATAATAAA 5

RESULT 9
LOCUS AR099868/c 29 bp DNA PAT 14-FEB-2001
DEFINITION Sequence 12 from patent US 6080400.
ACCESSION AR099868
VERSION AR099868.1 GI:12810316
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 29)
AUTHORS Williams,J.A. and Byrne,L.Marie.
TITLE Compositions for the prevention and treatment of verotoxin-induced disease
JOURNAL Patent: US 6080400-A 12 27-JUN-2000;
FEATURES Location/Qualifiers
SOURCE 1..29
BASE COUNT 11 a 2 c 5 g 11 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 29;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataataatttt 202
Db 22 AAATAATAATTTT 9

RESULT 10
LOCUS AR028182 30 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 31 from patent US 5858661.

```

ACCESSION   AR028182
VERSION     AR028182.1  GI:5940155
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 30)
AUTHORS     Shiloh,Y.
TITLE       Ataxia-telangiectasia gene and its genomic organization
JOURNAL     Patent: US 5858661-A 31-12-JAN-1999;
FEATURES
   source    1..30
              Location/Qualifiers
BASE COUNT   11 a      1 c      5 g      13 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 196 ttattttatttta 209
      |||||||
Db 14 TTAATTTTATTTTTA 27

RESULT 11
AR138585
LOCUS       AR138585 30 bp DNA
DEFINITION Sequence 110 from patent US 6200749.
ACCESSION   AR138585
VERSION     AR138585.1 GI:14480930
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 30)
AUTHORS     Shiloh,Y.
TITLE       Mutated forms of the ataxia-telangiectasia gene and method to
JOURNAL     Patent: US 6200749-A 110 13-MAR-2001;
FEATURES
   source    1..30
              Location/Qualifiers
BASE COUNT   11 a      1 c      5 g      13 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 196 ttattttatttta 209
      |||||||
Db 14 TTAATTTTATTTTTA 27

RESULT 12
E27913/c
LOCUS       E27913 32 bp DNA
DEFINITION Method for detecting foreign DNA fragment insert in Vero toxin
ACCESSION   E27913
VERSION     E27913.1 GI:13020766
KEYWORDS    JP 1999243996-A/3.
SOURCE      unidentified.
ORGANISM    unidentified.
REFERENCE   1 (bases 1 to 32)
AUTHORS     Masahiro,K.Y.N.N. and Kawamura,K.S.
TITLE       Method for detecting foreign DNA fragment insert in Vero toxin gene
JOURNAL     Patent: JP 1999243996-A 3 14-SEP-1999;
            TOYOCO CO LTD

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COMMENT      OS Unidentified
             PN JP 1999243996-A/3
             PD 14-SEP-1999
             PF 27-FEB-1998 JP 1998047677
             PR
             PI MASAHIRO KUSUMOTO,YOSHIKAKI NISHIYA,YOSHIHISA KAWAMURA, PI
             PC C1201/68,C12N15/09/(C12N15/09,C12R1:185),C12N15/00,
             (C12N15/00, PC C12R1:185)
             CC Strandedness: Both;
             CC Topology: Linear;
             CC Key
             FT source 1..32
              Location/Qualifiers
FEATURES
   source    1..32
              Location/Qualifiers
BASE COUNT   12 a      2 c      4 g      14 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 32;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 189 aaataattatttt 202
      |||||||
Db 17 AAAATATTATTATT 4

RESULT 13
AR019036/c
LOCUS       AR019036 36 bp DNA
DEFINITION Sequence 16 from patent US 5783196.
ACCESSION   AR019036
VERSION     AR019036.1 GI:3974150
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 36)
AUTHORS     Noriega,F.R. and Levine,M.M.
TITLE       Gua mutants of shigella spp. and vaccines containing the same
JOURNAL     Patent: US 5783196-A 16 21-JUL-1998;
FEATURES
   source    1..36
              Location/Qualifiers
BASE COUNT   11 a      3 c      10 g      12 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 189 aaataattatttt 202
      |||||||
Db 27 AAAATATTATTATT 14

RESULT 14
AR063204
LOCUS       AR063204 37 bp DNA
DEFINITION Sequence 8 from patent US 5844094.
ACCESSION   AR063204
VERSION     AR063204.1 GI:5990895
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 37)
AUTHORS     Hudson,P.John, Lah,M., Kortt,A.Andrew, Irving,R.Alexander,

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Atwell,J.Leslie, Malby,R.Louise, Power,B.Elaine and
Colman,P.Malcolm.
Target binding polypeptide
JOURNAL Patent: US 5844094-A 8 01-DEC-1998;
FEATURES Location/Qualifiers
source 1..37

BASE COUNT 14 a 5 c 0 g 18 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 37;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 83 aaattcataattat 96
Db 1 AAATTTATATATAT 14

RESULT 15
AX049973/c 45 bp DNA PAT 12-JAN-2001
LOCUS
DEFINITION Sequence 74 from Patent WO0070071.
ACCESSION AX049973
VERSION AX049973.1 GI:12226350
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Boul,A., Havenga,M.J. and Vogels,R.
TITLE Adenovirus derived gene delivery vehicles comprising at least one
JOURNAL element of adenovirus type 35
Patent: WO 0070071-A 74 23-NOV-2000;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Oligonucleotide TATA-plus"

BASE COUNT 14 a 5 c 7 g 19 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 79 ctgaaattataa 92
Db 21 CTGAAATTATATA 8

RESULT 16
AX049974 45 bp DNA PAT 12-JAN-2001
LOCUS
DEFINITION Sequence 75 from Patent WO0070071.
ACCESSION AX049974
VERSION AX049974.1 GI:12226351
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Boul,A., Havenga,M.J. and Vogels,R.
TITLE Adenovirus derived gene delivery vehicles comprising at least one
JOURNAL element of adenovirus type 35
Patent: WO 0070071-A 75 23-NOV-2000;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"

/db_xref="taxon:32630"
/note="Oligonucleotide TATA-m1n"
BASE COUNT 19 a 7 c 5 g 14 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 79 ctgaaattataa 92
Db 29 CTGAAATTATATA 42

RESULT 17
AX09885/c 45 bp DNA PAT 02-APR-2001
LOCUS
DEFINITION Sequence 4 from Patent WO0120014.
ACCESSION AX09885
VERSION AX09885.1 GI:13538881
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Schouten,G.J., Vogels,R. and Opstelten,D.J.
TITLE Modified adenoviral vectors for use in gene therapy
JOURNAL Patent: WO 0120014-A 4 22-MAR-2001;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Primer"
primer_bind 1..45
/note="Primer TATAp1us"

BASE COUNT 14 a 5 c 7 g 19 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 79 ctgaaattataa 92
Db 21 CTGAAATTATATA 8

RESULT 18
AX09886 45 bp DNA PAT 02-APR-2001
LOCUS
DEFINITION Sequence 5 from Patent WO0120014.
ACCESSION AX09886
VERSION AX09886.1 GI:13538882
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Schouten,G.J., Vogels,R. and Opstelten,D.J.
TITLE Modified adenoviral vectors for use in gene therapy
JOURNAL Patent: WO 0120014-A 5 22-MAR-2001;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Primer"
primer_bind 1..45
/note="Primer TATAp1us"

BASE COUNT 19 a 7 c 5 g 14 t

ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92
|||||
DB 29 CTGAATAATTATTA 42

RESULT 19

AXI37975/c 45 bp DNA PAT 30-MAY-2001
LOCUS AXI37975
DEFINITION Sequence 4 from Patent EP1083229.
ACCESSION AXI37975
VERSION AXI37975.1 GI:14274070
KEYWORDS

SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 45)

AUTHORS Modified adenoviral vectors for use in gene therapy
TITLE Patent: EP 1083229-A 4 14-MAR-2001;
JOURNAL Introgene B.V. (NL)

FEATURES
source location/Qualifiers

1.45
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="primer"

primer_bind 1.45
/note="Primer TATApIus"

BASE COUNT 14 a 5 c 7 g 19 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92
|||||
DB 21 CTGAATAATTATTA 8

RESULT 20

AXI37976 45 bp DNA PAT 30-MAY-2001
LOCUS AXI37976
DEFINITION Sequence 5 from Patent EP1083229.
ACCESSION AXI37976
VERSION AXI37976.1 GI:14274071
KEYWORDS

SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 45)

AUTHORS Modified adenoviral vectors for use in gene therapy
TITLE Patent: EP 1083229-A 5 14-MAR-2001;
JOURNAL Introgene B.V. (NL)

FEATURES
source location/Qualifiers

1.45
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="primer"

primer_bind 1.45
/note="Primer TATApIus"

BASE COUNT 19 a 7 c 5 g 14 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92
|||||
DB 29 CTGAATAATTATTA 42

RESULT 21

AXI59856/c 50 bp DNA PAT 22-JUN-2001
LOCUS AXI59856
DEFINITION Sequence 3184 from Patent WO0140521.
ACCESSION AXI59856
VERSION AXI59856.1 GI:14541187
KEYWORDS

SOURCE human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 50)

AUTHORS Shinkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and
methods of use thereof
JOURNAL Patent: WO 0140521-A 3184 07-JUN-2001;
Curagen Corporation (US)

FEATURES
source location/Qualifiers

1.50
/organism="Homo sapiens"
/db_xref="taxon:9606"
misc_feature 25..26
/note="Nucleotide deleted between bases 25 and 26
Accession number cg43063075"

misc_feature 26
/note="2 of 2 allelic variants (3183 is other entry)"
BASE COUNT 10 a 10 c 8 g 22 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 155 aaggaatatataa 168
|||||
DB 36 AAGGAATAATTATA 23

RESULT 22

ARI46953 18 bp DNA PAT 08-AUG-2001
LOCUS ARI46953
DEFINITION Sequence 13 from patent US 6221361.
ACCESSION ARI46953
VERSION ARI46953.1 GI:15110756
KEYWORDS

SOURCE unknown.
ORGANISM unknown.
REFERENCE 1 (bases 1 to 18)

AUTHORS Cochran,M.D. and Junker,D.E.
TITLE Recombinant swinepox virus
JOURNAL Patent: US 6221361-A 13 24-APR-2001;
FEATURES
source location/Qualifiers

1.18
/organism="unknown"
BASE COUNT 8 a 1 c 2 g 7 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgtaa 243
|||||
Db 5 AATGAATTGTAA 17

RESULT 23
LOCUS ARI21058 20 bp DNA
DEFINITION Sequence 79 from patent US 6159694.
ACCESSION ARI21058
VERSION ARI21058.1 GI:14104634
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Kairas,J.G.
TITLE Antisense modulation of stat3 expression
JOURNAL Patent: US 6159694-A 79 12-DEC-2000;
FEATURES
source location/Qualifiers
1..20
/organism="unknown"
BASE COUNT 9 a 3 c 3 g 5 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 173 attagataaag 185
|||||
Db 2 ATTGATATAAG 14

RESULT 24
LOCUS AX076045 20 bp DNA
DEFINITION Sequence 21 from Patent WO0104358.
ACCESSION AX076045
VERSION AX076045.1 GI:12710698
KEYWORDS
SOURCE Hepatitis B virus.
ORGANISM Hepatitis B virus.
REFERENCE 1 (bases 1 to 20)
AUTHORS Stuyver,L., Maertens,G. and van Geyl,C.
TITLE Detection of anti-hepatitis b drug resistance
JOURNAL Patent: WO 0104358-A 21 18-JAN-2001;
FEATURES
source location/Qualifiers
1..20
/organism="Hepatitis B virus"
/db_xref="taxon:10407"
BASE COUNT 6 a 3 c 4 g 7 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 72 tatatactgaaa 84
|||||
Db 15 TATATAGCTGAAA 3

RESULT 25
LOCUS A97479 23 bp DNA
DEFINITION Sequence 35 from Patent WO9916780.
ACCESSION A97479
VERSION A97479.1 GI:6780825

KEYWORDS unidentified.
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 23)
AUTHORS Gala,J. and Vannuffel,P.
TITLE GENETIC SEQUENCES, DIAGNOSTIC AND/OR QUANTIFICATION METHODS AND DEVICES FOR THE IDENTIFICATION OF STAPHYLOCOCCI STRAINS
JOURNAL Patent: WO 9916780-A 35 08-APR-1999;
FEATURES
source location/Qualifiers
1..23
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 11 a 2 c 3 g 7 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 23;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 58 aaagatatataa 70
|||||
Db 6 AAAGATATATATA 18

RESULT 26
LOCUS AX093544 24 bp DNA
DEFINITION Sequence 74 from Patent WO0118198.
ACCESSION AX093544
VERSION AX093544.1 GI:13509982
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 24)
AUTHORS Weissenbach,J. and Hazan,J.
TITLE Cloning, expression and characterisation of the spg4 gene responsible for the most frequent form of autosomal spastic paraplegia

JOURNAL Patent: WO 0118198-A 74 15-MAR-2001;
CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (FR)

FEATURES
source location/Qualifiers
1..24
/organism="Homo sapiens"
/db_xref="taxon:9606"
/note="Site accepteur d' passage du g ne spg4."

BASE COUNT 8 a 2 c 4 g 10 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 199 ttcttatlttaa 211
|||||
Db 3 TTTTATATTTTAAA 15

RESULT 27
LOCUS AX164353 24 bp DNA
DEFINITION Sequence 183 from Patent WO0138564.
ACCESSION AX164353
VERSION AX164353.1 GI:14545287
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequence.

REFERENCE 1 (bases 1 to 24)
AUTHORS Rouleau,G.A., Lafreniere,R.G., Rochefort,D., Cossette,P. and
Ragsdale,D.
TITLE Loci for idiopathic generalized epilepsy, mutations thereof and
method using same to assess, diagnose, prognosis or treat epilepsy
JOURNAL Patent: WO 0138564-A 183 31-MAY-2001;
McGill University (CA)
FEATURES
SOURCE 1..24
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="synthetic oligonucleotide"
BASE COUNT 2 a 5 c 3 g 14 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 301 ttatgtttcat 313
|||||
Db 8 TTTATGTTTCAAT 20

RESULT 28
AB069100 24 bp DNA SYN 08-AUG-2001
LOCUS AB069100/c
DEFINITION Synthetic construct DNA, forward primer for human STS-R133B1R
at 1p36.
ACCESSION AB069100
VERSION AB069100.1 GI:15129904
KEYWORDS
SOURCE synthetic construct DNA.
ORGANISM
REFERENCE 1 (bases 1 to 24)
AUTHORS Chen,Y.Z., Hayashi,Y., Wu,J.G., Takaoka,E., Maekawa,K.,
Watanabe,N., Inazawa,J., Hosoda,F., Arai,Y., Mizushima,H.,
Motohashi,A., Ohira,M., Nakagawara,A., Liu,S., Hoshii,M., Horii,A.
and Soeda,E.
TITLE A bac-based sts-content map spanning a 35-mb region of human
JOURNAL Chromosome 1p35-p36
MEDLINE 21269192
REFERENCE 2 (bases 1 to 24)
AUTHORS Horii,A.
TITLE Direct Submission
JOURNAL Submitted (04-AUG-2001) Akira Horii, Tohoku University School of
Medicine, Molecular Pathology; 2-1 Seiryomachi, Aoba-ku, Sendai,
Miyagi 980-8575, Japan (E-mail:horii@mail.cc.tohoku.ac.jp,
Tel:81-22-717-8042, Fax:81-22-717-8047)
FEATURES
SOURCE Location/Qualifiers
1..24
/organism="synthetic construct"
/db_xref="taxon:32630"
misc_feature 1..24
/note="forward primer for human STS-R133B1R at 1p36
STS-R133B1R obtained from clones B367G16, B133B1, B247E16,
B132L12, B32A16, B76C16, B81K9, B32I16, Human BAC library
RPC1-11"

BASE COUNT 9 a 2 c 4 g 9 t
ORIGIN

Query Match 4.0%; Score 13; DB 12; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 227 atataatgaatt 239
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Db 14 ATATAATGAATT 2

RESULT 29
AX042574/c
LOCUS AX042574 25 bp DNA PAT 23-NOV-2000
DEFINITION Sequence 140 from Patent WO0065088.
ACCESSION AX042574
VERSION AX042574.1 GI:11341182
KEYWORDS
SOURCE synthetic construct.
ORGANISM
REFERENCE 1 (bases 1 to 25)
AUTHORS Ulfendahl,P.J. and Wong,K.C.
TITLE Primers for identifying typing or classifying nucleic acids
JOURNAL Patent: WO 0065088-A 140 02-NOV-2000;
Amersham Pharmacia Biotech AB (SE)
FEATURES
SOURCE Location/Qualifiers
1..25
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="DQAI Homozygote primer sequence"
BASE COUNT 4 a 5 c 3 g 13 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 237 attgtaaaaaa 249
|||||
Db 17 ATTGTAATAAAAA 5

RESULT 30
AX043268/c
LOCUS AX043268 25 bp DNA PAT 23-NOV-2000
DEFINITION Sequence 834 from Patent WO0065088.
ACCESSION AX043268
VERSION AX043268.1 GI:11341876
KEYWORDS
SOURCE synthetic construct.
ORGANISM
REFERENCE 1 (bases 1 to 25)
AUTHORS Ulfendahl,P.J. and Wong,K.C.
TITLE Primers for identifying typing or classifying nucleic acids
JOURNAL Patent: WO 0065088-A 834 02-NOV-2000;
Amersham Pharmacia Biotech AB (SE)
FEATURES
SOURCE Location/Qualifiers
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/organism="synthetic construct"
/db_xref="taxon:32630"
/note="DQAI Heterozygote Primer Sequence"
BASE COUNT 4 a 5 c 3 g 13 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 237 attgtaaaaaa 249
|||||
Db 17 ATTGTAATAAAAA 5

RESULT 31
AX039624
LOCUS AX039624 26 bp DNA PAT 18-NOV-2000
DEFINITION Sequence 13 from Patent WO0063441.
ACCESSION AX039624
VERSION AX039624.1 GI:11229653

KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE
1 (bases 1 to 26)
AUTHORS Herrnstadt, C. and Davis, R. E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with alzheimer's disease
JOURNAL Patent: WO 0063441-A 13 26-OCT-2000;
MITOKOR (US)

FEATURES
source
1..26
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR primer"

BASE COUNT
8 a 2 c 5 g 11 t

ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 304 atgtttcatgtt 316
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DB 7 ATGTTTCATGTT 19

RESULT 32
AX039654 26 bp DNA PAT 18-NOV-2000
LOCUS AX039654
DEFINITION Sequence 43 from Patent WO0063441.
ACCESSION AX039654
VERSION AX039654.1 GI:11229683
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE
1 (bases 1 to 26)
AUTHORS Herrnstadt, C. and Davis, R. E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with alzheimer's disease
JOURNAL Patent: WO 0063441-A 43 26-OCT-2000;
MITOKOR (US)

FEATURES
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/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR primer"

BASE COUNT
8 a 2 c 5 g 11 t

ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 304 atgtttcatgtt 316
|||||

DB 7 ATGTTTCATGTT 19

RESULT 33
AR014028 28 bp DNA PAT 05-DEC-1998
LOCUS AR014028
DEFINITION Sequence 64 from patent US 5773246.
ACCESSION AR014028
VERSION AR014028.1 GI:3971482
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 28)
/organism="synthetic construct"
/db_xref="taxon:32630"

AUTHORS Keene, J.D., Levine, T. and Gao, F.
TITLE Methods and compositions useful in the recognition, binding and expression of ribonucleic acids involved in cell growth, neoplasia and immunoregulation
JOURNAL Patent: US 5773246-A 64 30-JUN-1998;
FEATURES
source
1..28
/organism="unknown"
BASE COUNT
6 a 3 c 1 g 18 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 196 ttattttatttt 208
|||||

DB 3 TTATTTTATTTT 15

RESULT 34
I21978 28 bp DNA PAT 07-OCT-1996
LOCUS I21978
DEFINITION Sequence 64 from patent US 5525495.
ACCESSION I21978
VERSION I21978.1 GI:1602332
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 28)
AUTHORS Keene, J.D., Levine, T. and Gao, F.
TITLE Methods and compositions useful in the recognition, binding and expression of ribonucleic acids involved in cell growth, neoplasia and immunoregulation
JOURNAL Patent: US 5525495-A 64 11-JUN-1996;
FEATURES
source
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/organism="unknown"
BASE COUNT
6 a 3 c 1 g 18 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 196 ttattttatttt 208
|||||

DB 3 TTATTTTATTTT 15

RESULT 35
AX012366 29 bp DNA PAT 06-SEP-2000
LOCUS AX012366
DEFINITION Sequence 26 from Patent EP0955369.
ACCESSION AX012366
VERSION AX012366.1 GI:9998412
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE
1 (bases 1 to 29)
AUTHORS Bartscher, H.D., Mueller, R.D., Hoelke, W.D. and Millan, J.L.
TITLE High active alkaline phosphatase
JOURNAL Patent: EP 0955369-A 26 10-NOV-1999;
ROCHE DIAGNOSTICS GMBH (DE)

FEATURES
source
1..29
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Artificial"

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BASE COUNT      8 a      7 c      9 g      5 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 29;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      22 gcacagaagaatg 34
      |||
      9 GCACAGAGAGATG 21

RESULT 36
E59972      29 bp      DNA      PAT      07-FEB-2001
LOCUS      Highly active alkaline phosphatase.
DEFINITION E59972
ACCESSION E59972
VERSION E59972.1 GI:13017742
KEYWORDS JP 199332586-A/23.
SOURCE JP 199332586-A/23.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 29)
AUTHORS Werner,H.R.M.M. and Burutoshu,J.L.M.M.
TITLE Highly active alkaline phosphatase
JOURNAL Patent: JP 199332586-A 23 07-DEC-1999;
          ROCHE DIAGNOSTICS GMBH
COMMENT OS Artificial Sequence
        PN JP 199332586-A/23
        PD 07-DEC-1999
        PE 06-MAY-1999 JP 1999126494
        PR 05-MAY-1998 DE 19819962;7
        PI WERNER HOERUKU,REINA MULLER,HERMUTTO BURUTOSHA, PI JOSE
        PC C12N15/09,C12N1/15,C12N1/19,C12N1/21,C12N5/10,C12N9/16, PC
        CC C12N15/00,C12N5/00
        FH Key
        FT source
        FT Location/Qualifiers
FEATURES
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    1..29      /organism="unidentified"
                /db_xref="taxon:32644"
BASE COUNT      8 a      7 c      9 g      5 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 29;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      22 gcacagaagaatg 34
      |||
      9 GCACAGAGAGATG 21

RESULT 37
AR028230      30 bp      DNA      PAT      29-SEP-1999
LOCUS      Sequence 79 from patent US 5858661.
DEFINITION AR028230
ACCESSION AR028230
VERSION AR028230.1 GI:5940203
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 30)
AUTHORS Shiloh,Y.
TITLE Ataxia-telangiectasia gene and its genomic organization
JOURNAL Patent: US 5858661-A 79 12-JAN-1999;
FEATURES
    Location/Qualifiers

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source      1..30
BASE COUNT      13 a      2 c      1 g      14 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 30;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      272 aaaaattatc 284
      |||
      15 AAAAATTATTC 27

RESULT 38
AR138633      30 bp      DNA      PAT      16-JUN-2001
LOCUS      AR138633
DEFINITION Sequence 158 from patent US 6200749.
ACCESSION AR138633
VERSION AR138633.1 GI:14480978
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 30)
AUTHORS Shiloh,Y.
TITLE Mutated forms of the ataxia-telangiectasia gene and method to
JOURNAL screen for a partial A-T phenotype
          Patent: US 6200749-A 158 13-MAR-2001;
          Location/Qualifiers
FEATURES
    source      Location/Qualifiers
    1..30      /organism="unknown"
BASE COUNT      13 a      2 c      1 g      14 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 30;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      272 aaaaattatc 284
      |||
      15 AAAAATTATTC 27

RESULT 39
AX063379      30 bp      DNA      PAT      24-JAN-2001
LOCUS      AX063379/C
DEFINITION Sequence 42 from Patent WO0079009.
ACCESSION AX063379
VERSION AX063379.1 GI:12541169
KEYWORDS
SOURCE
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 30)
AUTHORS Nazarenko,I. and Rashtchian,A.
TITLE Improved primers and methods for the detection and discrimination
JOURNAL of nucleic acids
          Patent: WO 0079009-A 42 28-DEC-2000;
          Life Technologies, Inc. (US)
FEATURES
    Location/Qualifiers
    1..30      /organism="synthetic construct"
                /db_xref="taxon:32630"
                /note="Oligonucleotide"
BASE COUNT      10 a      4 c      7 g      9 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 30;

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Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 40 atgaataaagat 52
|||||
Db 14 ATGAATAAAGAT 2

RESULT 40

195122/c

LOCUS 195122 31 bp DNA PAT .01-DEC-1998
DEFINITION Sequence 5 from patent US 5731425.
ACCESSION 195122
VERSION 195122.1 GI:3939592
KEYWORDS
SOURCE .
ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS 1 (bases 1 to 31)
Brizzard,B.L., Blanca,D.W., Chubet,R.G., Vizard,D.L. and
Hopp,T.Patrick.
TITLE Polypeptide surface marker for cells
JOURNAL Patent: US 5731425-A 5 24-MAR-1998;
FEATURES Location/Qualifiers
source 1. .31
BASE COUNT 12 a 11 c 7 g 1 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 31;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 124 gtgtcttcgagg 136
|||||
Db 28 GTGTCTCGGGG 16

RESULT 41

XELARSE59

LOCUS XELARSE59 33 bp DNA VRT 28-APR-1993
DEFINITION Xenopus laevis autonomous replication sequence e59.
ACCESSION R01606
VERSION R01606.1 GI:213953
KEYWORDS autonomous replication; mutational analysis.
SOURCE Xenopus laevis DNA.
ORGANISM Xenopus laevis
REFERENCE Unclassified.
AUTHORS Amphibia; Metazoa; Chordata; Vertebrata; Euteleostomi;
TITILE Amphibia; Batrachia; Anura; Mesobatrachia; Pipridae; Pipidae;
1 (bases 1 to 33)
Kearsey,S.
JOURNAL Structural requirements for the function of a yeast chromosomal
MEDLINE Cell 37, 299-307 (1984)
FEATURES 84205653
source 1. .33
BASE COUNT 8 a 4 c 2 g 19 t
ORIGIN /organism="Xenopus laevis"
/db_xref="taxon:8355"

Query Match 4.0%; Score 13; DB 5; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 298 attttatgttt 310
|||||
Db 1 ATTTTATGTTT 13

RESULT 42

A41027/c

LOCUS A41027 36 bp DNA PAT 05-MAR-1997
DEFINITION Sequence 14 from Patent EP0626449.
ACCESSION A41027
VERSION A41027.1 GI:2296916
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified
REFERENCE Unclassified.
AUTHORS 1 (bases 1 to 36)
Schreier,P.H., Stenzel,K.D., Adam,G.P. and Maiss,E.D.
TITLE DNA coding for plant virus sequences
JOURNAL Patent: EP 0626449-A 14 30-NOV-1994;
BAYER AG (DE)
COMMENT Other publication JP 6343469 941220
Other publication CA 2124272 941129
Other publication CN 1098744 950215
Other publication AU 6191494 941201
Other publication DE 4317845 941201
Other publication ZA 9403730 950202.
FEATURES Location/Qualifiers
source 1. .36
BASE COUNT 9 a 5 c 5 g 17 t
ORIGIN /organism="unidentified"
/db_xref="taxon:32644"

Query Match 4.0%; Score 13; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 156 aggaataataaa 168
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Db 24 AGGAATAATATAA 12

RESULT 43

AR082586/c

LOCUS AR082586 36 bp DNA PAT 31-AUG-2000
DEFINITION Sequence 14 from patent US 5973135.
ACCESSION AR082586
VERSION AR082586.1 GI:10009308
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS 1 (bases 1 to 36)
Schreier,P.Helmolt, Stenzel,K., Adam,G. and Maiss,E.
TITLE DNA comprising plum pox virus and tomato spotted wilt virus CDNAS
for disease resistance
JOURNAL Patent: US 5973135-A 14 26-OCT-1999;
FEATURES Location/Qualifiers
source 1. .36
BASE COUNT 9 a 5 c 5 g 17 t
ORIGIN /organism="unknown"
/db_xref="taxon:32644"

Query Match 4.0%; Score 13; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 156 aggaataataaa 168
|||||
Db 24 AGGAATAATATAA 12

RESULT 44

AX167671

LOCUS AX167671 36 bp DNA PAT 03-JUL-2001
DEFINITION Sequence 16 from Patent WO0144277.

ACCESSION AX167671 GI:14597058
 VERSION AX167671.1

KEYWORDS
 SOURCE synthetic construct.
 ORGANISM synthetic construct.
 artificial sequence.

REFERENCE 1 (bases 1 to 36)

AUTHORS Weglich Glover, L., Budziszewski, G. J., Levin, J. Z., and Zhou, Q.

TITLE Herbicide target genes and methods

JOURNAL Patent: WO 0144277-A 16 21-JUN-2001;

SYNGENTA PARTICIPATIONS AG (CH)

FEATURES
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/organism="synthetic construct"

/db_xref="taxon:32630"

/note="Oligonucleotide"

BASE COUNT 6 a 4 c 11 g 15 t

ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 36;

Best Local Similarity 100.0%; Pred. No. 1.6e+05; Mismatches 0; Gaps 0;

Matches 13; Conservative 0; Indels 0;

QY 303 tatgtttcatgt 315

DB 21 TATGTTTTCATGT 33

RESULT 45

LOCUS 128261 36 bp DNA

DEFINITION Sequence 14 from patent US 5569823. PAT 06-FEB-1997

ACCESSION 128261

VERSION 128261.1 GI:1819037

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 36)

AUTHORS Schreier, P. H., Stenzel, K., Adam, G., Unter and Maiss, E.

TITLE DNA comprising plum pox virus and tomato spotted wilt virus CDNAS

JOURNAL Patent: US 5569823-A 14 29-OCT-1996;

FEATURES location/Qualifiers

source 1..36

/organism="unknown"

BASE COUNT 9 a 5 c 5 g 17 t

ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 36;

Best Local Similarity 100.0%; Pred. No. 1.6e+05; Mismatches 0; Gaps 0;

Matches 13; Conservative 0; Indels 0;

QY 156 aggaataataa 168

DB 24 AGGAATAATATAA 12

Search completed: January 24, 2002, 03:22:36
 Job time: 3813 sec

SUMMARIES

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1	327	100.0	327	6	AX004614	AX004614 Sequence
2	327	100.0	1392	1	L77965	L77965 Clostridium
3	327	100.0	1392	6	AX004613	AX004613 Sequence
4	199	60.9	54310	1	AP003515	AP003515 Clostridi
5	81.8	25.0	12900	3	AE001429	AE001429 plasmodu
6	79	24.2	39347	9	AL135966	AL135966 Human DNA
7	77.8	23.8	53932	2	AC023371	AC023371 Homo sapi
8	77.4	23.7	163443	2	AC006280	AC006280 plasmodu
9	77.4	23.7	205449	2	AC005506	AC005506 plasmodu
C 10	76.2	23.3	3392	3	AF300334	AF300334 Dictyoste
C 11	75.8	23.2	242513	2	AC079314	AC079314 Homo sapi
C 12	75.8	23.2	318231	2	PFMA113P3	PFMA113P3 plasmodu
C 13	75.6	23.1	140414	2	AF377947	AF377947 Oryza sat
C 14	75	22.9	178783	2	AC068139	AC068139 Homo sapi
C 15	74.4	22.8	156060	2	AC004153	AC004153 plasmodu
C 16	73.2	22.4	863	11	CNS06EVC	AL395628 T7 endo c
17	73	22.3	8622	8	YSCMTCYTCOC	M97514 Saccharomyc
C 18	72.8	22.3	34119	8	AF3222718	AF3222718 Chrysidid
C 19	72.2	22.1	104992	2	AC005504	AC005504 plasmodu
20	72.2	22.1	162445	2	AL158151	AL158151 Human DNA
21	72.2	22.1	169546	2	AC004157	AC004157 plasmodu
22	72.2	22.1	199882	2	AL354720	AL354720 Human DNA
23	72	22.0	158398	2	AC011146	AC011146 Homo sapi
C 24	71.6	21.9	180934	9	AE0073409	AE0073409 Homo sapi
C 25	71.4	21.8	12029	3	ABE001400	ABE001400 plasmodu
C 26	71.4	21.8	175053	2	AC090014	AC090014 Homo sapi
C 27	71.2	21.8	110000	2	AL591074_2	Continuatiuon (3 of
C 28	71	21.7	95477	9	AC007076	AC007076 Homo sapi
C 29	71	21.7	168799	9	AC009531	AC009531 Homo sapi
C 30	71	21.7	194038	9	AC010103	AC010103 Homo sapi
C 31	70.8	21.7	159475	2	AC021378	AC021378 Homo sapi
C 32	70.6	21.6	13433	3	AF315648	AF315648 Ceratilis
C 33	70.6	21.6	85779	8	SC6011856	AJ011856 Saccharom
C 34	70.6	21.6	122747	2	AC093220	AC093220 Homo sapi
C 35	70.6	21.6	159255	2	AF212831	AF212831 Homo sapi
36	70.6	21.6	161230	2	AC011355	AC011355 Homo sapi
37	70.6	21.6	234112	3	PFMA14P2	AL035475 plasmodu
38	70.6	21.6	340000	9	HS21C013	AL163213 Homo sapi
C 39	70.4	21.5	155456	2	AC027753	AC027753 Homo sapi
C 40	70.4	21.5	160624	2	AC060835	AC060835 Homo sapi
C 41	70.4	21.5	172758	2	AC022553	AC022553 Homo sapi
C 42	70.4	21.5	199551	2	AC006281	AC006281 plasmodu
C 43	70.2	21.5	137342	9	AL392048	AL392048 Human DNA
C 44	70.2	21.5	180388	9	H0MRERBLAS	L19150 Human reclin
C 45	70.2	21.5	183584	9	AC012492	AC012492 Homo sapi

BASE COUNT 141 a 13 c 44 g 129 t
ORIGIN

Query Match 100.0%; Score 327; DB 6; Length 327;

Best Local Similarity 100.0%; Pred. No. 3.5e-28; Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 attggatattcttaaatctagcagagaagaatgtttaaatgaataataaaaaa 60
1 ATTTGGATATCTTAAATTTAGCACAGAAATGTTTAAATGAATTAAGATTAATAA 60
61 gatataattatatactgaataattataatataatgataatgataatgaataa 120
61 GATATATTAATTAATATAGCTGAAAATTTATTAATTAATGATAGTATGTTAATAA 120
121 aaagtgtctcgaggagacacttttctttaaagaagaataataataaattagat 180
121 AAAGGTCTCGGGGACACTTTTGTGTTTAAAGAAATTAATTAATTAATTAATGAT 180
181 aaagtgttaataatt 240
181 AAAAGTGAATTAATTAATTTTAAATTTGTTTAAATTTGATTAATTAATTAATG 240
241 taaaaaaattcgaaggagaaatacaatgaataaattatcgaagtttactgaat 300
241 TAAAAAAATTTCAAGGGGGAATTAATGAATAAATTTATTTCAAGTTTACTGTAAT 300
301 ttatgtttcattgtttcttcttattgtt 327
301 TTATGTTTTCATGTTTCTTATTTGTT 327

RESULT 2

LOCUS L77965 1392 bp DNA BCT 28-JUL-1998
DEFINITION Clostridium perfringens C beta 2 toxin gene, complete cds.
ACCESSION L77965
VERSION L77965.1 GI:3342214
KEYWORDS
SOURCE Clostridium perfringens C.
ORGANISM Clostridium perfringens C.
Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae; Clostridium.

REFERENCE 1 (bases 1 to 1392)
AUTHORS Gilbert, M., Jolivet-Reynaud, C. and Popoff, M. R.
TITLE Beta2 toxin, a novel toxin produced by Clostridium perfringens
JOURNAL Gene 203 (1), 65-73 (1997)
MEDLINE 98085977
ERRATUM: [[published erratum appears in Gene 1998 Mar 27;210(1):173]]
2 (bases 1 to 1392)
REFERENCE Popoff, M. R.
AUTHORS Direct Submission
TITLE Submitted (15-JAN-1998) Toxines Microbiennes, Institut Pasteur,
JOURNAL Paris cedex 15 75724, France
COMMENT GSDS:S:76036.
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location retrieved from GSDS Fri Jul 24 15:39:17 1998].

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source
1. .1392
/organism="Clostridium perfringens C"
/strain="CWC245"
/db_xref="taxon:79668"
255..260
/note="putative"
RBS 268..357
/note="putative"
sig_peptide 268..1065
/note="putative"
CDS /codon_start=1
/transl_table=1
/product="beta 2 toxin"

/protein_id="AAC27654.1"
/db_xref="GI:3342215"
/translation="MKRTISKFTYIFMFSCLYIGALISPMKASAKKEIDATRKVENVTL
NALKNYDINTVAVNISSEDERVNNVQREYREMLDFKYPDPQQLKSEIILNSOKSDNKEIF
NKTTEFLNGALYIDMEFTVSSKDKGLIYSDMERTRVENCKYILTPSPFTQYCTWDEL
AOAIGEVYAPQYSDREFTYADNIIILNFRQVATSGSRDLKVEYSVVDHMMKDDVKASQ
MVQNDPSARQILYLTIEKGOSFYKRYRIRINLFPASIRVGEBCA"
358..1062
/note="putative"
terminator /product="beta 2 toxin"
1066..1104
/note="putative"

BASE COUNT 606 a 115 c 209 g 462 t
ORIGIN

Query Match 100.0%; Score 327; DB 1; Length 1392;
Best Local Similarity 100.0%; Pred. No. 2.8e-28; Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 attggatattcttaaatctagcagagaagaatgtttaaatgaataataaaaaa 60
1 ATTTGGATATCTTAAATTTAGCACAGAAATGTTTAAATGAATTAAGATTAATAA 60
61 gatataattatatactgaataattataatataatgataatgataatgaataa 120
61 GATATATTAATTAATATAGCTGAAAATTTATTAATTAATGATAGTATGTTAATAA 120
121 aaagtgtctcgaggagacacttttctttaaagaagaataataataaattagat 180
121 AAAGGTCTCGGGGACACTTTTGTGTTTAAAGAAATTAATTAATTAATTAATGAT 180
181 aaagtgttaataatt 240
181 AAAAGTGAATTAATTAATTTTAAATTTGTTTAAATTTGATTAATTAATTAATG 240
241 taaaaaaattcgaaggagaaatacaatgaataaattatcgaagtttactgaat 300
241 TAAAAAAATTTCAAGGGGGAATTAATGAATAAATTTATTTCAAGTTTACTGTAAT 300
301 ttatgtttcattgtttcttcttattgtt 327
301 TTATGTTTTCATGTTTCTTATTTGTT 327

RESULT 3

LOCUS AX004613 1392 bp DNA PAT 24-AUG-2000
DEFINITION Sequence 1 from Patent WO9915669.
ACCESSION AX004613
VERSION AX004613.1 GI:9928053
KEYWORDS
SOURCE Clostridium perfringens.
ORGANISM Clostridium perfringens.
Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae; Clostridium.

REFERENCE 1 (bases 1 to 1392)
AUTHORS Gilbert, M. and Popoff, M. R.
TITLE Clostridium toxin, and method for preparing immunogenic
JOURNAL Compositions
PATENT: WO 9915669-A 1 01-APR-1999;
PATENT: WO 9915669-A 1 01-APR-1999;
JOURNAL GIBERT MARYSE (FR); PASTEUR INSTITUT (FR)

FEATURES
source
1. .1392
/organism="Clostridium perfringens"
/db_xref="taxon:1502"
268..1065
/note="unnamed protein product"
CDS /codon_start=1
/transl_table=1
/protein_id="CAC04901.1"
/db_xref="GI:9928054"
/translation="MKRTISKFTYIFMFSCLYIGALISPMKASAKKEIDATRKVENVTL

NALNKNDINTVNISEDERNVNNQYREMLEDEKYPDNQOLKSEFILLNSOKDNKEIF
 NVKEFGNAGIYDEFTVSXKDGKLIYSDEMRKVENEGKYLTPSPRTQVCMWDEL
 AOAATGYPQYDREYFYADNILLNFRQATSGSRDLKVEYSVVDMMKDKVKASQ
 MVIQONPDSRQRLRIETEKQOSYKRIKRIKNEFPASIRVFGGYCA

BASE COUNT 606 a 115 c 209 g 462 t

ORIGIN

Query Match 100.0%; Score 327; DB 6; Length 1392;
 Best Local Similarity 100.0%; Pired. No. 2.8e-28;
 Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 attgggagatccttaaatagcacaagaagtgttaaatgaataaagaataa 60
 Db 1 ATTGGGAGATCTTAAATTAGCACAGAAATGTTAAATGAATAAAGATAA 60
 Qy 61 gatataatataatagctgaanaattataatataatgaatagtaataa 120
 Db 61 GATATATAATATATAGCTGAATAATTATATATAGTATAGTAAATAA 120
 Qy 121 aaagtgcttcggggggaacttttctttaaagaagaataataaattagat 180
 Db 121 AAAGTGCTTCGGGGGACCTTTTGTTTAAAGGAAATATATAAATTTAGAT 180
 Qy 181 aaagtgaataataatttttatttaattgttaaaaaattgataatgaattg 240
 Db 181 AAAAGTGAATAATATTTTATTATTTTAAATTTGTTAAATTTGATATATGAATTG 240
 Qy 241 taaaaaaaattcaggggggaataataatgaaaaaattattcaaaattactgtaatt 300
 Db 241 TAAAAAAATTTACGGGGGAGATATAATGAAAAAATTTATCAAGTTACTGTAAT 300
 Qy 301 ttatgttttcattgttttctatctgt 327
 Db 301 TTATGTTTTCATGTTTCTTATGTT 327

RESULT 4
 AP003515 54310 bp DNA circular BCT 10-AUG-2001
 DEFINITION Clostridium perfringens plasmid pCP13 DNA, complete sequence.
 ACCESSION AP003515
 VERSION AP003515.1 GI:15076712
 KEYWORDS
 SOURCE Clostridium perfringens (strain:13) plasmid:PCP13 DNA.
 ORGANISM Clostridium perfringens
 Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;

REFERENCE 1 (bases 1 to 54310)
 AUTHORS Ohtani,K., Ohshima,S., Hirakawa,H., Ohshima,K., Shida,T.,
 Shimizu,T., Hattori,M., Kuhara,S., Hayashi,H. and Shimizu,T.
 TITLE Complete Nucleotide Sequence of the Virulence Plasmid pCP13 from
 Clostridium perfringens

JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 54310)
 AUTHORS Shimizu,T.
 TITLE Direct Submission
 JOURNAL Submitted (12-APR-2001) Tohru Shimizu, Institute of Basic Medical
 Sciences, University of Tsukuba, Department of Microbiology; 1-1-1
 Tennoudai, Tsukuba, Ibaraki 305-8575, Japan
 E-mail:tschimizuend.tsukuba.ac.jp, Tel:81-298-53-3354,
 Fax:81-298-53-3354)

FEATURES
 source Location/Qualifiers
 1..54310
 /organism="Clostridium perfringens"
 /plasmid="pCP13"
 /strain="13"
 /db_xref="taxon:1502"
 /note="anaerobic pathogen for gas gangrene"
 gene 940..1692
 /gene="soj"
 CDS 940..1692
 /gene="soj"

/note="250 aa, similar to p1r:14044 Sp00A activation
 inhibitor soj from Bacillus subtilis (253 aa); 378
 identity in 250 aa overlap
 pCP01

Para family"
 /codon_start=1
 /transl_table=11
 /product="Soj protein"
 /protein_id="BAB62438.1"
 /db_xref="GI:15076713"
 /translation="MKRISVFNKIGVAKTTSTANGACLEKGRVLLVLDPOSNL
 TKLFKAVSMEDVASIADVLIDKNDIEKVIKTFDENIDILPSVTLAFARKILLDVN
 RSOONRLAKLEETEDKDYDCLIDCPALMTITVNALCASDEVLPVPIKIDKFDLIGLE
 YLDSIEETKDEFPNPNIEKGFITMDSSSTTVKVKYIKQLKSVLGKEMRTSIHONIK
 VVESTIECPVVS SKKASLWKDLSKEIF"
 1751..3031
 /gene="parB"
 1751..3031
 /gene="parB"
 /note="426 aa, similar to gpu:AF300944_3 presumptive ParB
 protein from Lactococcus lactis subsplactis (242 aa); 308
 identity in 266 aa overlap
 pCP02"

/codon_start=1
 /transl_table=11
 /product="ParB protein"
 /protein_id="BAB62439.1"
 /db_xref="GI:15076714"
 /translation="MAKFSISGCMNGISKNTKRYEERQAKENKFEYINIDIRKN
 EKNFEIVLDESIAEDIKLGNLNVRLNDMDLELISGERRYALSKLVENKKE
 KNIYPPCKVIESNDIDSEIILIOANQSRLEVEKLTVOERLOEYKIKKENGKVP
 KIRDIIDLNDLSATQVGRYERINNKILPEIKAVITOGNTTANASEPSSISENRY
 ILSIIDKTRMSQEAVDLKNKIKIIEKEKELELKKAYEKELELRLBEKKNOYK
 LKSENEKLRKLDNNIIEBKETBGQIIEFEKLNKENVILIEELSKYDKKIEDI
 TKAEKNNEKQRLKDELKSLKRSNNEVDIKTKENFVLVONLKLIDNSFKMKSQI
 NKKKENVAVAEETKAKEPLEKYKEISDLKTL"
 3147..3509
 /gene="pCP03"
 3147..3509
 /gene="pCP03"
 /note="120 aa, no significant homology"
 /codon_start=1
 /transl_table=11
 /product="hypothetical protein"
 /protein_id="BAB62440.1"
 /db_xref="GI:15076715"
 /translation="MEKILAEKRINISFYKRRKGAALVTLIYLPKMLEVIGITENERE
 CFFYIEDKAIKISKEKQSEAKERTISFSTSYKTYLNNKMLEYLGVSDEDSCTIEL
 RKKDITLVKRDNGRIDIDI"
 3773..4024
 /gene="pCP04"
 3773..4024
 /gene="pCP04"
 /note="83 aa, similar to p1r:T14710 probable transposase
 from Yersinia pestis (402 aa); 44% identity in 50 aa
 overlap
 truncated"

gene 4040..4222
 /gene="pCP05"
 4040..4222
 /gene="pCP05"
 /note="60 aa, similar to gp:AF143819.1 transposase-like
 protein from Escherichia coli (402 aa); 38% identity in 60
 aa overlap
 truncated"
 /codon_start=1

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gene
CDS
/translation="MKNRKLKIFETIASLLVGSIGTTALATVTHSSDKFEASLPGLFS
/protein_id="BAB62443.1"
/db_xref="GI:15076718"
/translation="MISKHQRNIRNIEDKNLNLVASCITTRDVAGQIKALYDIETSAE
TVSNITNRIMPLVSEW"
4588..4746
/gene="PCP06"
4588..4746
/gene="PCP06"
/note="52 aa, similar to p1r:T43600 probable transposase
from Yersinia pestis (105 aa); 56% identity in 50 aa
overlap
truncated"
/codon_start=1
/translation="1"
/product="probable transposase"
/protein_id="BAB62444.1"
/db_xref="GI:15076719"
/translation="MLYTNVIESLNSQFRKFTKTLFIPNDVSLIKMLYATEKVNK
KWTNRYNR"
complement(5169..5804)
/gene="PCP07"
complement(5169..5804)
/gene="PCP07"
/function="ATP-binding protein"
/note="211 aa, similar to gp:AE001508.4 ABC transporter
(ATP-binding protein) from Bacillus halodurans (213 aa);
49% identity in 214 aa overlap"
/codon_start=1
/translation="1"
/product="probable ABC transporter"
/protein_id="BAB62445.1"
/db_xref="GI:15076720"
/translation="MNTIEISNLKKRYFDKIFKDFSLISIKGEMIAISGSGCKST
LNLMLGLEKFDSEIITDGVKNIKNSKLNKFLREKISTYLFQNFALVDEYVEENL
RLAIKHTIKNTKRIIEEIIICLKEFVGLGCKKNYIELSGEQORVALARMLKPEEI
ILADEPTGSLDENRDIISLKEINESGKTIITVTHDNYAKADRIIFL"
complement(5804..7966)
/gene="PCP08"
complement(5804..7966)
/gene="PCP08"
/note="720 aa, similar to gp:AE001508.3 BH0280 gene
product from Bacillus halodurans (713 aa); 23% identity in
661 aa overlap"
/codon_start=1
/translation="1"
/product="conserved hypothetical protein"
/protein_id="BAB62446.1"
/db_xref="GI:15076721"
/translation="MKKKVALIILFIITLISFPGVYSVRNHTEFMKLNLOCNENF
EVTISIDREYNAYAESITKSLEYNINIFSEVDINEDYRKYINCYFENKEENH
LPIVSGRFRHKNDISTYITDSEDTOQIVINDFNKNLHEITIKSKDINTEFN
LFIQIENEQILDKLIEDLKSESIIVQKVGSDSQYETFLKILLVCFGLFMIF
YQVIGSYKIKIGIKLGHSTFVMLKERLEVLREIVLMLVTVLFIENFKNTSLF
WKFMELTICISIMIEFIVSIVPIYVSKITSLNINKRPVKSIIINSPVAVIL
ASLIFPSNALDLSISIGKEKNYKWEETKOYIILPEIGPNDESTOSPISEMEK
ERAYIVYKOGATILADPNRYEPRSMKAKOMLPEYVRETIIVNPNLKHKYVDV
GNINISDEKDRILVPEKTRNEKELLETLYGINSQPCSTTCSHTAGCKMLVE
QOKRITIMKSNQKYFSLDVNPEGNVTPPIVSLFESDKLVSYKIIIGYNNSPF
KIRANSEEVINGLEKYYDMGVILDPNLDVNASTIINIRAKVKVIFPIVILLAV
ISIILOMTSLYFNQNNKRIIVKRLHGYRLIRYKNYFIMVLITWTCPLAASLITKD
INIITYLIVLIVELVFIFENINSLEKKNLKVIGKEV"
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/gene="PCP09"
complement(8043..8354)
/gene="PCP09"
/note="103 aa, no significant homology"
/codon_start=1
/translation="1"
/product="hypothetical protein"
/protein_id="BAB62447.1"
/db_xref="GI:15076722"
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gene
CDS
/translation="MKRNLKIFETIASLLVGSIGTTALATVTHSSDKFEASLPGLFS
QAQSSKFFCGEQKRRATARVAVGTTLVEAKNIKAKILAHNQCTYYYGVEBMNSYVNH
L"
8779..9012
/gene="PCP10"
8779..9012
/gene="PCP10"
/note="77 aa, similar to probable transposase from
Yersinia pestis plasmid pMT1 (402 aa); 25% identity in 158
aa overlap
truncated"
/codon_start=1
/translation="1"
/product="probable transposase"
/protein_id="BAB62448.1"
/db_xref="GI:15076723"
/translation="MKELVSLICMSTNEGSKFMLSPPHKKEFAKDLTIYGSVNETG
MKNDLEIREKWSKYPNVVKSWMKDNMDNLSTFF"
9127..9366
/gene="PCP11"

Query Match      60.9%; Score 199; DB 1; Length 54310;
Best Local Similarity 86.3%; Pred. No. 2.2e-14;
Matches 220; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

QY 73 atatgctgaaattatataattatgataagtgatgaataaataaagtgcttcg 132
DB 13459 ATGAAATTTTAACTCATGCTTTTAAAGTTAATATATATTTTAAATTTAGGTGCCAC 13518
QY 133 ggggacacttttgctttaaaggaaataataaataattagataaaggctgtaa 192
DB 13519 GGGGACATCTTTTGTGTTTAAAGTAAATATGATTAATTTAGATGATAAAAGTAA 13578
QY 193 taattatttattttaaattgttaaattgataatgaattgtaaaaaaatt 252
DB 13579 GAATTAATTTTAAATTTTAAATTTGTTAAATTTGATATATGAAATTTGAAAAAATTT 13638
QY 253 caggggggataataataaataattatcaagttactgtaattttagtttca 312
DB 13639 CAGGGGGGATATTAATGAAAAAATTTATTCAAAGTTTACTGTAATTTTATGTTTCA 13698
QY 313 tgtttcttattgtt 327
DB 13699 TATTTTCTTATGTT 13713

RESULT 5
AE001429 12900 bp DNA INV 06-NOV-1998
LOCUS Plasmodium falciparum chromosome 2, section 66 of 73 of the
DEFINITION complete sequence.
ACCESSION AE001429 AE001362
VERSION AE001429.1 GI:3845321
KEYWORDS
SOURCE
ORGANISM
Plasmodium falciparum.
REFERENCE
1 (bases 1 to 12900)
Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
Gardner,M.J., Tettein,H., Carucci,D.J., Cummings,L.M., Aravind,L.,
Koonin,E.V., Shalim,S., Mason,T., Yu,K., Fujii,C., Pederson,J.,
Shen,K., Jing,J., Aston,C., Lai,D., Schwartz,D.C., Perdeson,M.,
Salzberg,S., Zhou,L., Sutton,G.G., Clayton,R., White,O.,
Smith,H.O., Fraser,C.M., Hoffman,S.L., et.al.
Chromosome 2 sequence of the human malaria parasite Plasmodium
falciparum
Science 282 (5391), 1126-1132 (1998)
JOURNAL MEDLINE 99021743
REMARK Erratum: [[published erratum appears in Science 1998 Dec
4;282(5395):1827]]
2 (bases 1 to 12900)
REFERENCE
Gardner,M.J.
Direct Submission
JOURNAL Submitted (02-NOV-1998) The Institute for Genomic Research, 9712
```

Medical Center Drive, Rockville, MD 20814, USA

FEATURES
Source

1.12900
/organism="Plasmodium falciparum"
/db_xref="taxon:5833"
/chromosome="2"
11045..12083
/gene="PF0955W"
join(11045..11113,11229..12083)
/gene="PF0955W"
/note="Identified by sequence similarity; putative"

CDS
/product="rflin"
/codon_start=1
/protein_id="AAC71980.1"
/db_xref="GI:3845322"

/translation="MNTYIMLMVMSILLVLEISYVNNHNKYNNGYIONNFQIM
KLSRRLEIQLPKCPHYNDPELKIIDKLEERIKRYIEFNNSFEELHGLVERK
SLYNGKSSNMEKELIKKYDSDIRDEHNVISGSIYSDYRLVYAKSEYONKIL
RDELASCCVNDYLDNLKKGCFGSGVIGICLSLVNSNIGYIVLAKSEYITL
DIANKFTKLAGIYFFSSSIENAGSGVITFYWDSMRMASIASSTINPYGIALVLI
VLVYLVLYVIMVTRRRKSKMKHCKHLS"

BASE COUNT 5356 a 845 c 1167 g 5532 t
ORIGIN

Query Match 25.0%; Score 81.8; DB 3; Length 12900;

Best Local Similarity 54.0%; Pred. No. 0.25; Mismatches 142; Indels 0; Gaps 0;

Matches 167; Conservative 0; Mismatches 142; Indels 0; Gaps 0;

FEATURES

Source

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TrEMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr6> RPI-20N11 is from the library RPI-1 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/> VECTOR: pcypac2

IMPORTANT: This sequence is not the entire insert of clone RPI-20N11. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.

The true right end of clone RPI-20N11 is at 3947 in this sequence. The true right end of clone RPI-48213 is at 103 in this sequence.

Location/Qualifiers

1..39347
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="6"
/clone="RPI-20N11"
/clone_11b="RPI-1"

repeat_region

/note="L1P1A12 repeat: matches 1421..-291 of consensus" 3460..3893

repeat_region /note="L1M3C repeat: matches 999..1402 of consensus" 3894..4213

repeat_region /note="AluY repeat: matches 1..311 of consensus" 4214..4415

repeat_region /note="L1M3C repeat: matches 1402..1610 of consensus" 4426..4777

repeat_region /note="MER47A repeat: matches 13..366 of consensus" 4779..5831

repeat_region /note="L1M2 repeat: matches 1528..2663 of consensus" 6028..6073

repeat_region /note="L1 repeat: matches 2729..2777 of consensus" 6091..8167

repeat_region /note="L1P1A15 repeat: matches 4085..6157 of consensus" 8287..8567

repeat_region /note="AluX repeat: matches 3..284 of consensus" 8792..8965

repeat_region /note="L1 repeat: matches 2737..2901 of consensus" 8959..9103

repeat_region /note="L1M3C repeat: matches 1092..1236 of consensus" 9227..10104

repeat_region /note="L1 repeat: matches 2903..3922 of consensus" 10091..10657

repeat_region /note="L1M2 repeat: matches 5115..5687 of consensus" 10657..11436

repeat_region /note="L1P1A2 repeat: matches 3..776 of consensus" 11432..16672

repeat_region /note="L1P1A2 repeat: matches 900..6146 of consensus" 16682..16795

repeat_region /note="L1P3 repeat: matches 5677..5791 of consensus" 16821..16860

repeat_region /note="20 copies 2 mer ta 90% conserved" 16865..17035

COMMENT : On Mar 15, 2000 this sequence version replaced gi:7210154.

RESULT 6
AL135906 39347 bp DNA PRI 26-APR-2000
LOCUS Human DNA sequence from clone RPI-20N11 on chromosome 6 Contains
DEFINITION GSSS, complete sequence.
ACCESSION AL135906
VERSION AL135906.19 GI:7248200
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 39347)
AUTHORS Blakey,S.
TITLE Direct Submission
JOURNAL Submitted (14-APR-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
Requests: clonerequest@sanger.ac.uk
On Mar 15, 2000 this sequence version replaced gi:7210154.

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repeat_region      17036. .17343
                    /note="MER4 repeat: matches 367. .646 of consensus"
repeat_region      17377. .17678
                    /note="LIM49 repeat: matches 5783. .6112 of consensus"
repeat_region      17679. .17996
                    /note="ALUyB8 repeat: matches 1. .318 of consensus"
repeat_region      17997. .18193
                    /note="LIM49 repeat: matches 6112. .6300 of consensus"
repeat_region      19452. .20219
                    /note="LIME repeat: matches 429. .1193 of consensus"
misc_feature       complement(20068. .20528)
                    /note="match: GSS: Em:AQ218253"
repeat_region      23091. .23131
                    /note="MLTII repeat: matches 370. .409 of consensus"
repeat_region      23144. .23391
                    /note="LIP44 repeat: matches 5999. .6146 of consensus"
repeat_region      25530. .25553
                    /note="L12 copies 2 mer ca 95% conserved"
repeat_region      25570. .25754
                    /note="ALUto/FRAM repeat: matches 173. .291 of consensus"
repeat_region      25827. .26622
                    /note="398 copies 2 mer aa 58% conserved"
misc_feature       25879. .26639
                    /note="Tandem repeat. Tandem repeat contains forced join.
                    BamII and EcoRI bot suggest approx 200bp missing."
misc_feature       complement(26230. .26730)
                    /note="match: GSS: Em:B69019"
misc_feature       complement(26267. .26664)
                    /note="match: GSS: Em:AQ230058"
repeat_region      27019. .27243
                    /note="LTR40a repeat: matches 282. .516 of consensus"
repeat_region      27306. .28098
                    /note="LIME2 repeat: matches 5094. .5922 of consensus"
repeat_region      28440. .28705
                    /note="L13 copies 2 mer aa 54% conserved"
repeat_region      29274. .29834
                    /note="LIMB6 repeat: matches 5552. .6128 of consensus"
repeat_region      29885. .31038
                    /note="LIM4 repeat: matches 862. .2044 of consensus"
repeat_region      31742. .32332
                    /note="LIMD1 repeat: matches 5317. .5893 of consensus"
misc_feature       complement(32119. .32398)
                    /note="match: GSS: Em:AQ078965"
repeat_region      32769. .32872
                    /note="52 copies 2 mer tt 64% conserved"
misc_feature       complement(32852. .33057)
                    /note="match: GSS: Em:AQ780759"
repeat_region      33046. .33127
                    /note="41 copies 2 mer tt 73% conserved"
repeat_region      33143. .34362
                    /note="LIP47 repeat: matches 4855. .6127 of consensus"
repeat_region      34564. .34625
                    /note="31 copies 2 mer ta 71% conserved"
repeat_region      34637. .34664
                    /note="14 copies 2 mer ac 89% conserved"
repeat_region      34745. .36725
                    /note="LIP13 repeat: matches 2266. .4198 of consensus"
repeat_region      36726. .37032
                    /note="ALU repeat: matches 3. .308 of consensus"
repeat_region      37033. .38914
                    /note="LIP13 repeat: matches 4198. .6145 of consensus"
repeat_region      39090. .39341
                    /note="HIGHER2 repeat: matches 2448. .2717 of consensus"
BASE COUNT      14429 a 7155 c 6738 g 11025 t
ORIGIN

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Query Match      24.2% Score 79; DB 9; Length 39347;
Best Local Similarity 54.7% Pred. No. 0.43;
Matches 157; Conservative 0; Mismatches 130; Indels 0; Gaps 0;
Qy 24 acagaagaatggttaataaagataaagataaataataataataagctgaa 83
   | | | | | | | | | | | | | | | | | | | | | | | | | | | |

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Db 25934 ATAAATATATATAAAATATATTATATATATAAAATATATATAAAAT 25993
Qy 84 aattataattatagataagatagttcaataaataaaggtctcgggggaactt 143
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Db 25994 AATATATATATATAAAATATATAAAATATATATAAAATATATATAAAAT 26053
Qy 144 ttgttttaaaaaaggaatacaataaattagataaagtcgaataattttt 203
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Db 26054 ATATATAAATATATATAAAATATATATATATATAAAATATATATAAATATATATAT 26113
Qy 204 atttcaaatcttcaaaaattgacataactgaatctgaaaaaatttcagggggaat 263
   | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 26114 ATAAATATATATATAAAATATATATATATATAAAATATATATAAATATATATAAATAT 26173
Qy 264 ataataagaaaaaattatttcaagttacgttaattttttagttt 310
   | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 26174 AATATATATATATAAAATATATATAAAATATATATAAATATATATAT 26220

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RESULT 7
AC023371      53932 bp      DNA      HTG      13-JUL-2000
DEFINITION      Homo sapiens clone RP11-21D18, 10M-PASS SEQUENCE SAMPLING.
ACCESSION      AC023371
VERSION      AC023371.2 GI:9123990
KEYWORDS      HTG; HTGS_PHASE0.
SOURCE      human.
ORGANISM      Homo sapiens
                Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

```

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REFERENCE
AUTHORS      1 (bases 1 to 53932)
                Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE      Homo sapiens, clone RP11-21D18
JOURNAL      Unpublished
REFERENCE
AUTHORS      2 (bases 1 to 53932)
                Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
                Anderson,S., Baldwin,J., Barna,N., Beda,F., Boguslavsky,L.,
                Bouckgalter,B., Brown,A., Burkett,G., Campoliano,A., Castle,A.,
                Choquet,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,
                Dearellano,K., Dewar,K., Dodge,S., Domino,M., Doyle,M.,
                Feneator,J., Ferreira,P., Fitzhugh,M., Forrest,C., Gage,D.,
                Galagan,J., Gardyna,S., Glnde,S., Goyette,M., Graham,L.,
                Grand-pierre,N., Grant,G., Hagos,B., Heatford,A., Horton,L.,
                Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
                Klein,J., Landers,T., Larcocque,R., Lehoczy,J., Levine,R.,
                Lieu,C., Liu,G., Locke,K., MacDonald,P., Marquis,N., McCarthy,M.,
                McKern,P., McGurt,A., McKernan,K., McPheters,R., Meldrum,J.,
                Meneus,L., Milnova,T., Miranda,C., Mienga,V., Morrow,J., Naylor,J.,
                Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Olivat,T.M.,
                Peterson,K., Pierre,N., Pisanl,C., Pollara,V., Raymond,C.,
                Riley,R., Rogov,P., Rothman,D., Roy,A., Santos,R., Schauer,S.,
                Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
                Subramanian,A., Talamas,J., Testife,S., Theodore,J., Tirrell,A.,
                Travers,M., Trigglio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B.,
                Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zimmer,A. and
                Zody,M.

```

```

TITLE      Direct Submission
JOURNAL
COMMENT      Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome
                Research, 320 Charles Street, Cambridge, MA 02141, USA
                On Jul 13, 2000 this sequence version replaced gi.6970502.
                All repeats were identified using RepeatMasker:
                Smit, A.F.A. & Green, P. (1996-1997)
                http://ftp.genome.washington.edu/RM/RepeatMasker.html
                ----- Genome Center

```

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Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: L3985
Center clone name: 21_D18

```

* NOTE: This record contains 55 individual

[illegible]


```

Sequence Data
JOURNAL      Unpublished
REFERENCE    2 (bases 1 to 178783)
AUTHORS      Smith,D.R.
TITLE        Direct Submission
JOURNAL      Submitted (29-Apr-2000) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
REFERENCE    3 (bases 1 to 178783)
AUTHORS      Smith,D.R.
TITLE        Direct Submission
JOURNAL      Submitted (15-Nov-2000) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
COMMENT      On Nov 15, 2000 this sequence version replaced gi:9887668.
FEATURES
  source
    1..178783
    /organism="Homo sapiens"
    /db_xref="taxon:9606"
    /chromosome="10"
    /clone="RP11-506M13"
    /clone_id="RPC1-11"
BASE COUNT   53204 a 37175 c 38013 g 50391 t
ORIGIN
Query Match      22.9% Score 75; DB 9; Length 178783;
Best Local Similarity 53.6%; Pred. No. 0.93;
Matches 156; Conservative 0; Mismatches 135; Indels 0; Gaps 0;
QY 14 taaattgacagagaagaattgtaaatgaaataaagaataaagaataataa 73
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 150164 TAAATATATTATATATATATATATATATATATATATATATATAT 150105
QY 74 tatagctgaattataattatagatagatagatgtaataaagaagcttcg 133
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 150104 TATATATATATATATATATATATATATATATATATATATATAT 150045
QY 134 gggacacttttctgttaaaaggaataataaataaattgataaagtcgtaaat 193
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 150044 ATTATATATATATATATATATATATATATATATATATATATAT 149985
QY 194 aactatttataaatttgtaaaattgataaattgtaattgtaaaaaatttc 253
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 149984 TATATATATATATATATATATATATATATATATATATATATAT 149925
QY 254 agggggggaattataaataaataaattatcaagttcgttaatttta 304
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 149924 ATAAATATATATATATATATATATATATATATATATATATAT 149874

RESULT 15
AC004153 156060 bp DNA HTG 12-AUG-2000
LOCUS      Plasmodium falciparum chromosome 12 clone 3D7, *** SEQUENCING IN
DEFINITION PROGRESS **, 2 unordered pieces.
ACCESSION AC004153
VERSION    AC004153.7 GI:9797733
KEYWORDS   HTG; HTGS; PHASE1.
SOURCE     malaria parasite P. falciparum.
ORGANISM   Plasmodium falciparum
            Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
REFERENCE 1 (bases 1 to 156060)
AUTHORS    Hyman,R.W., Qin,F., Fung,E.L., Conway,A.B. and Davis,R.W.
TITLE      Kurdi,O.B., Conway,A.B. and Davis,R.W.
JOURNAL    Plasmodium falciparum 3D7 chromosome 12
            Unpublished
            2 (bases 1 to 156060)
REFERENCE 2 (bases 1 to 156060)
AUTHORS    Hyman,R.W., Qin,F., Fung,E.L., Conway,A.B. and Davis,R.W.
TITLE      Direct Submission
JOURNAL    Submitted (18-FEB-1998) Stanford DNA Sequencing and Technology
            Center, Stanford University, 855 California Avenue, Palo Alto, CA
            94304, USA
COMMENT     On Aug 12, 2000 this sequence version replaced gi:8810454.
            * NOTE: This is a 'working draft' sequence. It currently

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* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 102169: contig of 102169 bp in length
* 102170 102369: gap of unknown length
* 102370 156060: contig of 53691 bp in length.
FEATURES
  source
    1..156060
    /organism="Plasmodium falciparum"
    /db_xref="taxon:5833"
    /chromosome="12"
    /clone="PFYAC812"
    /clone_id="3D7"
BASE COUNT   62615 a 14889 c 15137 g 63219 t 200 others
ORIGIN
Query Match      22.8% Score 74.4; DB 2; Length 156060;
Best Local Similarity 54.9%; Pred. No. 1.1;
Matches 167; Conservative 0; Mismatches 136; Indels 1; Gaps 1;
QY 24 acagaaagatgtttaaatgaaataaagaataaagaataaataaataa 83
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 31682 AAATTAATAAATAAATAAATAAATAAATAAATAAATAAATAAATA 31741
QY 84 aattataattatagataagatagtaataaataaagaagtcctcgaggacatt 143
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 31742 AAATTAATAAATAAATAAATAAATAAATAAATAAATAAATAAATA 31800
QY 144 ttgttttaaaaggaataataaataaattgataaagaagtcgtaaatatttt 203
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 31801 AATTAAAAAATAAATAAATAAATAAATAAATAAATAAATAAATA 31860
QY 204 atttaatttgtaaaattcgataaattgtaattgtaaaaaaattcaaggagggaat 263
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 31861 TTTTTCATTATATATATATATATATATATATATATATATATAT 31920
QY 264 ataaatgaaaaaattatttcaaggttaccgtaattttatglttccatttctat 323
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 31921 AACATTAATAAATAAATAAATAAATAAATAAATAAATAAATAAATA 31980
QY 324 tggc 327
    ||
DB 31981 ATTT 31984

```

Search completed: January 24, 2002, 02:24:45
 Job time: 438 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:10:08 ; Search time 2272.52 Seconds
(without alignments)
1546.244 Million cell updates/sec

Title: US-09-531-438-3

Perfect score: 327
Sequence: 1 atttggaatacttaattt.....ttcatgtttctattgtt 327

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 11351937 seqs, 5372889281 residues

Total number of hits satisfying chosen parameters: 22703874

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database :

EST:*
1: em_estfun:*
2: em_esthum:*
3: em_estin:*
4: em_estom:*
5: em_estpl:*
6: em_estba:*
7: em_estro:*
8: em_estov:*
9: em_hic:*
10: gb_estl:*
11: gb_est2:*
12: gb_hic:*
13: gb_gss:*
14: em_gss_fun:*
15: em_gss_hum:*
16: em_gss_inv:*
17: em_gss_pln:*
18: em_gss_pro:*
19: em_gss_rdg:*
20: em_gss_vrt:*
21: em_gss_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	91	27.8	1101	13	CNS00EVL
2	89.2	27.3	1225	13	CNS0161D
3	85.4	26.1	1101	13	CNS003BD
4	84.6	25.9	1101	13	CNS0039G
5	84.4	25.8	905	13	CNS00KHK
6	83.2	25.4	524	13	CNS01U90
7	83	25.4	1101	13	CNS008X3
8	83	25.4	1101	13	CNS00E07
9	82.8	25.3	987	13	CNS014P0
10	82.2	25.1	1101	13	CNS0014P
11	81.2	24.8	734	13	CNS010MP
12	81.2	24.8	1200	13	CNS016C0

C 13	81	24.8	1125	10	AL547503
C 14	80.6	24.6	1092	13	CNS020K7
C 15	80.4	24.6	973	13	CNS0080F
C 16	80.4	24.6	1001	13	CNS0155H
C 17	80.4	24.6	1101	13	CNS00FMC
C 18	80.2	24.5	836	13	CNS02W02
C 19	79.6	24.3	694	13	AO853360
C 20	79.4	24.3	1101	13	CNS00EVL
C 21	79	24.2	966	13	CNS0052C
C 22	79	24.2	1101	13	CNS0021J
C 23	78.6	24.0	581	13	CNS034DK
C 24	78.6	24.0	928	13	CNS00DXY
C 25	78.6	24.0	996	13	CNS00FJH
C 26	78.4	24.0	1069	13	CNS0107G
C 27	78.4	24.0	1101	13	CNS00E07
C 28	78.4	24.0	1101	13	CNS016L1
C 29	78	23.9	876	13	CNS00961
C 30	77.8	23.8	987	13	CNS014P0
C 31	77.6	23.7	1101	13	CNS00FYG
C 32	77.6	23.7	1101	13	CNS00238
C 33	77.4	23.7	770	13	AO740708
C 34	77.4	23.7	1101	13	CNS000B8
C 35	77.4	23.7	1225	13	CNS00EPO
C 36	77.4	23.7	1101	13	CNS0161D
C 37	77.2	23.6	1101	13	CNS00KAE
C 38	77.2	23.6	1200	13	CNS016C0
C 39	76.8	23.5	614	13	CNS0152H
C 40	76.8	23.5	1099	10	AL536986
C 41	76.6	23.4	1101	13	CNS0039Q
C 42	76.4	23.4	1092	13	CNS020K7
C 43	76.2	23.3	928	13	CNS00DKY
C 44	76.2	23.3	1203	13	CNS015MU
C 45	76	23.2	1124	13	CNS073BM

ALIGNMENTS

RESULT 1
CNS00EVL 1101 bp DNA GSS 04-JUN-1999
LOCUS Drosophila melanogaster genome survey sequence T7 end of BAC:
DEFINITION BACR29B23 of Rpci-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE
AUTHORS
TITLE
JOURNAL

COMMENT

Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see <http://www.fruitfly.org> The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osoegawa and
Aaron Mammose in Pieter de Jong's laboratory in the Department of
Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,
NY. The library is named Rpci-98 and was constructed by partial
EcoRI digestion of Drosophila DNA provided by the BDGP from the
isogenic strain y2; cn bw sp, the same strain used for the BDGP's
PI and EST libraries. A more detailed description of the library
and how to order individual BAC clones, the entire library, or
filters for hybridization from the BACPAC Resource Center can be
found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

```

FEATURES
source
1. 1101
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone_11b="RPC1-98"
/clone="BACR29B23"
/note="end : 17"

BASE COUNT      419 a      91 c      60 g      299 t      232 others

ORIGIN

Query Match      27.8%  Score 91; DB 13; Length 1101;
Best Local Similarity 41.6%  Pred. No. 0.0093;
Matches 127; Conservative 68; Mismatches 108; Indels 2; Gaps 1;

Oy 15 aaattgacacgaagaatgtttaatgaataaagaataaagaataaagaataatcattat 74
||| : : : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 700 AAAAAAAMATWAAWAAATWATWAAATWATWAAATWATWAAATWATWAAATWATWAAATW
75 atagctgaaattcataatcataagataagtagtaataataaagaagtcctcg9 134
||| : : : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 760 ATATATATWTTWAAWAAATWAAWAAATWATWAAATWATWAAATWAAWAAATWATWAAATW
Oy 135 ggaacacttttgtc--ttaaagaagaaataataaataattcgataaagtcgtaaa 192
||| : : : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 820 AAAMWAAWATWAAWAAATWATWAAATWAAWAAATWAAWAAATWAAATWAAATWATWAAWAA
193 taattatttttaatttaatttgcgttaaaatctgataatcgaattgataaagaat 252
||| : : : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 880 TAAWATTTWTTTATWAAWAAATWATWAAATWAAWAAATWAAWAAATWAAWAAATWATWAAW
Oy 253 caaggagggaataataaagaaataattcctaagttactcgtaattttatgtttca 312
||| : : : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 940 WATATTWTTATTAATWAAATWATWAAATWATWAAATWATWAAATWATWAAATWATWAAW
Oy 313 tggtt 317
||| : : : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 1000 TAAWAT 1004

RESULT 2
LOCUS CNS0161D/c
DEFINITION Drosophila melanogaster genome survey sequence Sp6 end of BAC
BACN1518 of DrosBAC library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
ACCESSION AL106171
VERSION AL106171.1 GI:5620504
KEYWORDS GSS.
SOURCE fruit fly
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 1225)
Genoscope.
Direct Submission
Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : sequef@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
determination of this BAC-end sequence was carried out as part of a
collaboration with the European Drosophila Genome Project (EDGP) -
http://www.edgp.ebi.ac.uk/. This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billard at CPH (Centre
d'Etude du Polymorphisme Humain) with funding provided by a MRC
project grant. The DNA was prepared from embryos by Alain Bucheton
and Genevieve Payan. It has been constructed in the vector
pbeloBAC11.

FEATURES
source
1. 1225
/organism="Drosophila melanogaster"
/plasmid="pbeloBAC11"
/db_xref="taxon:7227"

```

BASE COUNT	266 a	128 c	38 g	368 t	425 others
ORIGIN					
Query Match	27.3%	Score 89.2:	DB 13:	Length 1225:	
Best Local Similarity	40.8%	Pred. No. 0.014:			
Matches 125:	Conservative 61:	Mismatches 120:	Indels 0:	Gaps 0:	
OY	8	atattcctaatttagcacagaagaatggtttaaattgaataaagaataaataaagatatat 67			
Db	1164	ATATAMATMTATTTATTTWMAAAAMMTTTTATATATATMTATWATAATAAAATAATATAMW 1105			
OY	68	taattatatagctgaaaaatttataatttatatgataagcatgtaattcaataaaaaagcgt 127			
Db	1104	WAMWMAAAAMWMAAAAMWMAWMTWMAAAATATATAATATATATAATAATWMAWMAW 1045			
OY	128	tctcg9ggacacttttctgttttaaaaggaaaaataaataaattagataaaagctg 187			
Db	1044	TWMAAAATATATATTTTTTTTTTTTNAAAAAAAMAAATAATATATWATAATAAAMW 985			
OY	168	taaaataatttatttattcaattgltcaaaaattgataataatgaaattgtaaaaaa 247			
Db	984	TAAAAAAAAATAAATATTATWMAWMTTTTTTAAAMWMTTTTTTTTTWMTTAAAMWT 925			
OY	248	aatttcagg9ggaaataaataaataaataattattcaagtttaccgttaattttatgt 307			
Db	924	WMTTATMTTATTWTRRRARAATWMTTTTTTTTTTTTTTTTATATTTTTTTTMMHT 865			
OY	308	tttcat 313			
Db	864	TYTMT 859			
RESULT 3					
CNS003BD/c					
LOCUS	CNS003BD	1101 bp	DNA	GSS	03-JUN-1999
DEFINITION	Drosophila melanogaster genome survey sequence TE73 end of BAC # BACR08K08 of RPCI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.				
ACCESSION	AI064091	1	GI:4941847		
VERSION	AI064091.1				
KEYWORDS	GSS.				
ORGANISM	fruit fly.				
SOURCE	Drosophila melanogaster				
REFERENCE	Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.				
AUTHORS	I (bases 1 to 1101)				
TITLE	Genoscope.				
JOURNAL	Direct Submission				
COMMENT	Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : seque@genoscope.cns.fr - Web : www.genoscope.cns.fr)				
FEATURES	Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see http://www.fruitfly.org The BDGP Drosophila melanogaster BAC library was prepared by Kazuoto Osoegawa and Aaron Mammeter in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's pl and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm .				
SOURCE	Location/Qualifiers				
	1..1101				

Submitted 22-JUN-1999) Genoscope - Centre National de Séquençage BP 101 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr - Web : www.genoscope.cns.fr)

Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the *Drosophila melanogaster* genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP *Drosophila melanogaster* BAC library was prepared by Kazutoyo Oseegawa and Aaron Mammeter in Pierer de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPci-98 and was constructed by partial EcoRI digestion of *Drosophila* DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's p1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or

filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

source

1. 905
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone.lib="RPCI-98"
/clone="BACR17N06"
/note="end : 17"

BASE COUNT 388 a 82 c 77 g 194 t 164 others
ORIGIN

Query Match

Best Local Similarity 25.8%; Score 84.4; DB 13; Length 905;
Pred No. 0.063; Mismatches 129; Indels 1; Gaps 1;

Matches 141; Conservative 55; Mismatches 129; Indels 1; Gaps 1;

Qy 2 ttgggatacttaattagcagagaagatgtttaaataagaataaagaag 61
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 581 DTKMKWKTAMKAAAAAATAATWATATWATATAAATAATATAATWAAAAA 640
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 62 atataataatataagcgaataatataatataatgaatgaatgaat 121
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 641 AAAAAAATAAATAATATWTTTAAATAAATAAATAAATAAATAAATAA 700
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 122 aagtgctcgggagacattttgttttaaaagaataataaataatga 181
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 701 AAAAAAATAAATAATWTTTAAATAAATAAATAAATAAATAAATAA 760
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 182 aagtgtaataataatattttattttaattgttaaaattgataatga 241
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 761 AAW-ARWGATATATATATAAATAATWATATWATATWATATWATATA 819
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 242 aaaaaaatttcagggggaataataaataaataatttcaaaagttac 301
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 820 AAWATTAATAAATAATATAAATAAATAAATAAATAAATAAATAATTT 879
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 302 ttatgtttcattgtttcttattgtt 327
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 880 TTTTWT 905

RESULT 6

CNS01090/c

LOCUS 524 bp DNA GSS 12-MAY-2000
DEFINITION Tetradon nigroviridis genome survey sequence PUC-Orl end of clone
196c24 of library G from Tetradon nigroviridis, genomic survey
sequence.

ACCESSION

AL167541
AL167541.1 GI:7805598

KEYWORDS GSS: genome survey sequence.
SOURCE Tetradon nigroviridis.

ORGANISM

Tetradon nigroviridis.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontidae; Tetradon.

REFERENCE

1 (bases 1 to 524)
Roest-Crollius, H., Jallou, O., Dasilva, C., Fizames, C., Fisher, C.,
Bouneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and
Weissenbach, J.

TITLE

Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetradon nigroviridis

JOURNAL

2 (bases 1 to 524)
Roest-Crollius, H., Jallou, O., Dasilva, C., Bouneau, L., Fisher, C.,
Bernot, A., Fizames, C., Wincker, P., Brothier, P., Quetier, F.,
Saurin, W. and Weissenbach, J.

TITLE

Human gene number estimate provided by genome wide analysis using
Tetradon nigroviridis DNA sequence

JOURNAL

3 (bases 1 to 524)
Genoscope.

TITLE Direct Submission

JOURNAL

Submitted (12-APR-2000) to the EMBL/Genbank/DBSI databases
This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetradon nigroviridis
genome. For more information, please take a look at
<http://www.genoscope.cns.fr/Tetradon>.

COMMENT

FEATURES

source

1. 524
/organism="Tetradon nigroviridis"
/db_xref="taxon:99883"
/clone.lib="196C24"
/clone="196C24"
/note="Genoscope sequence ID : C0AG196B12SP1-end :
PUC-Orl"

BASE COUNT 124 a 15 c 24 g 298 t 63 others
ORIGIN

Query Match

Best Local Similarity 25.4%; Score 83.2; DB 13; Length 524;
Pred No. 0.12; Mismatches 129; Indels 0; Gaps 0;

Matches 133; Conservative 46; Mismatches 129; Indels 0; Gaps 0;

Qy 15 aaattagcacagaagatgtttaaataagaataaagaatataat 74
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 399 AAAAAAATAAATAATWATATWATATAAATAAATAAATAAATAAATA 340
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 75 atagcgaataattataatataatgaatgaatgaatgaatgaatga 134
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 339 TTTAAATTTATTTATTTATTTATTTATTTATTTATTTATTTATTT 280
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 135 ggaacattttgtttaaaagaataataaataaattgataaagtcga 194
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 279 AAAAAAATAAATAAATAAATAAATAAATAAATAAATAAATAAATA 220
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 195 attatttatttataattgtttaaaattgataaattgataaattca 254
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 219 TATATTATTATTTATTTATTTATTTATTTATTTATTTATTTATTT 160
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 255 gggggaataataaataaataaataaatttcaagttactgtaattt 314
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 159 AAAATTAATAAATAAATAAATAAATAAATAAATAAATAAATA 100
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Qy 315 ttcttcta 322
| : : : : : | : : : : : | : : : : : | : : : : : | : : : : :
Db 99 TTTTMMWW 92

RESULT 7

CNS008X3

LOCUS 1101 bp DNA GSS 03-JUN-1999
DEFINITION Drosophila melanogaster genome survey sequence T7 end of BAC #
BACR18L14 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.

ACCESSION

AL052544
AL052544.1 GI:4934295

KEYWORDS

GSS:
GSS.

SOURCE

fruit fly.
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

1 (bases 1 to 1101)
Direct Submission
Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)
- Web : www.genoscope.cns.fr

JOURNAL

Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see <http://www.fruitfly.org> The BDGP Drosophila
melanogaster BAC library was prepared by Kazuo Osoegawa and

COMMENT

Aaron Mammoser in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

Source

Location/Qualifiers

1. 1101

/organism="Drosophila melanogaster"

/db_xref="taxon:7227"

/clone_lib="RPCI-98"

/clone="BACR18L14"

/note="end : 17"

BASE COUNT 251 a 151 c 169 g 239 t 291 others

ORIGIN

Query Match 25.4%; Score 83; DB 13; Length 1101;

Best Local Similarity 37.3%; Pred. No. 0.082;

Matches 113; Conservative 74; Mismatches 116; Indels 0; Gaps 0;

Oy 13 ttaaatgacagcaagaatcgttaaatgaataaagataataaagataataatt 72

Db 799 DDAKAAATAAAGAAADADAAAARAKAAATKAAAGAAKATTAAGKRAAGAAK 858

Oy 73 atatagctgaataattcaataatataatgaatagatagataaagaagcttcg 132

Db 859 TTTTAAWAAWAAATTTTAAWTTTAAWAAWAAWAAWAAWAAWAAWAAWAAW 918

Oy 133 ggggagcacttttctttaaagaagaataataaattcgaataaagctaa 192

Db 919 ARAKAAATATWMAATATAADTATAAATAATATATATATATATATATATATAT 978

Oy 193 taattatttattttaaatttgaataatttgaataatttgaataatttgaataatt 252

Db 979 ATRMAATTTTAT 1038

Oy 253 caggggggaataataataataattcgaagttcgaattcgaattcgaattcga 312

Db 1039 WMAAATAAATAAT 1098

Oy 313 tgt 315

Db 1099 TWT 1101

RESULT 8

CNS00E07

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

COMMENT

COMMENT

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COMMENT

please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazuhiro Osoegawa and Aaron Mammoser in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

Source

Location/Qualifiers

1. 1101

/organism="Drosophila melanogaster"

/db_xref="taxon:7227"

/clone_lib="RPCI-98"

/clone="BACR29P01"

/note="end : 173"

BASE COUNT 366 a 66 c 104 g 351 t 214 others

ORIGIN

Query Match 25.4%; Score 83; DB 13; Length 1101;

Best Local Similarity 45.0%; Pred. No. 0.082;

Matches 144; Conservative 48; Mismatches 119; Indels 9; Gaps 1;

Oy 8 atatcttaatttgcagcaagaatcgttaaatgaataaagataataaagataat 67

Db 557 ATKTTTATATWMAATATATAAATAATTTTAAATTTTAAATTTTAAATTTTAA 616

Oy 68 taattatagctgaataattcaataatataatgaatagatagataaagaagct 127

Db 617 AAWTAT 676

Oy 128 tctcgggggacacttttctttaaagaagaataataaattcgaataaagct 187

Db 677 TA-----AMAWTTTAAWTTATTAARWMAATTTAAWMAAAWTTTAAWMAAAT 727

Oy 188 taaataatttatttattttaaatttgaataatttgaataatttgaataatttga 247

Db 728 AAAATATAATTTTATTTTAAWMAWTTTAAATTTTAAWMAWTTTAAWMAWTT 787

Oy 248 aattcaggggggaataataataataattcgaagttcgaattcgaattcga 307

Db 788 ATNWTATATAATTTTAAWTTTAAWTTTAAWTTTAAWTTTAAWTTTAAWTT 847

Oy 308 ttcatglttcttatttgt 327

Db 848 WTTTAAWTTTAAWTTTATTTT 867

RESULT 9

CNS014PQ/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

<http://www.edgp.ebi.ac.uk> - This Drosophila melanogaster BAC library (Dros BAC) was made by Alain Billand at CEPH (Centre d'Etude du Polymorphisme Humain) with funding provided by a MRC project grant. The DNA was prepared from embryos by Alain Bucheton and Genevieve Payan. It has been constructed in the vector pBelobAC11.

FEATURES

source

Location/Qualifiers
1..987
/organism="Drosophila melanogaster"
/plasmid="pBelobAC11"
/db_xref="taxon:7227"
/clone_lib="DrosBAC"
/clone="BACN12P22"
/note="end : SP6"

BASE COUNT 257 a 122 c 122 g 241 t 245 others
ORIGIN

Query Match 25.3%; Score 82.8; DB 13; Length 987;
Best Local Similarity 37.0%; Pred. NO. 0.093;

Matches 118; Conservative 79; Mismatches 121; Indels 1; Gaps 1;

QY 8 atactctaaattagcacagaagaattglttaaatgaataaagataaagaatataat 67
DB 952 WWWWWWWWWTTTTTTTTTTTTTTTTTTTTTAAWWTTTAAWWWWWWWWTTTTTT 893
QY 68 taattatagctgaataattataatataagtagtataataataaagtg 127
DB 892 TTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTT 833
QY 128 tctcgaggacacatttcttcttaaaaagaataataaataaataaagtg 187
DB 832 WNTNTAAWMTTATTAATTTAAWMTTAAWMTTAAWMTTAAWMTTAAWMTTAA 774
QY 188 taaaataatttatttatttatttatttatttatttatttatttatttattt 247
DB 773 TTAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTT 714
QY 248 aatttcagaggaggaataataaataaataaataaataaataaataaataa 307
DB 713 AAAMAMMAM 654
QY 308 ttcatgtttcttcttcttcttcttcttcttcttcttcttcttcttctt 326
DB 653 TMMTKMTTK 635

RESULT 10

CNS001FB/C

LOCUS

DEFINITION

CNS001FB 1101 bp DNA GSS 03-JUN-1999
Drosophila melanogaster genome survey sequence TET3 end of BAC # BACR04A23 of RPCT-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

fruit fly.
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 1101)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Direct Submission
Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqrefgenoscope.cns.fr
- Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see <http://www.fruitfly.org> The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osoegawa and

Aaron Mammosser in Pieter de Jong's laboratory in the Department of
Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,
NY. The library is named RPCT-98 and was constructed by partial
EcoRI digestion of Drosophila DNA provided by the BDGP from the
isogenic strain y2; cn bw sp, the same strain used for the BDGP's
P1 and EST libraries. A more detailed description of the library
and how to order individual BAC clones, the entire library, or
filters for hybridization from the BACPAC Resource Center can be
found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

source

Location/Qualifiers
1..1101
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone_lib="RPCT-98"
/clone="BACR04A23"
/note="end : TET3"

BASE COUNT 288 a 110 c 103 g 491 t 109 others
ORIGIN

Query Match 25.1%; Score 82.2; DB 13; Length 1101;
Best Local Similarity 43.4%; Pred. NO. 0.1;

Matches 135; Conservative 42; Mismatches 134; Indels 0; Gaps 0;

QY 17 attgacacagaagaattglttaaatgaataaagaataaagaatataatataat 76
DB 1101 AT 1042
QY 77 agctgaataattataactatataatgaatagtagtgaataaagaagtcgcgggg 136
DB 1041 ATAAAAAAAT 982
QY 137 acacatttcttcttaaaaagaataataaataaataaataaataaataa 196
DB 981 AT 922
QY 197 tatttatttataatttcttgaataattgataataattgataaataaattcagg 256
DB 921 ATTAAMMTAT 862
QY 257 gagggaataaataaataaataaataaataaataaataaataaataaataa 316
DB 861 AAAAMNTAT 802
QY 317 tctctattgt 327
DB 801 AT 791

RESULT 11

CNS010MP/C

LOCUS

DEFINITION

CNS010MP 734 bp DNA GSS 26-JUL-1999
Drosophila melanogaster genome survey sequence T7 end of BAC BACN04L20 of DrosBAC library from Drosophila melanogaster (fruit fly), genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

fruit fly.
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 734)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Direct Submission
Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqrefgenoscope.cns.fr
- Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a
collaboration with the European Drosophila Genome Project (EDGP) -
<http://www.edgp.ebi.ac.uk> - This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billand at CEPH (Centre

	http://fulllength.invitrogen.com"				
BASE COUNT	415 a	142 c	148 g	345 t	75 others
ORIGIN					

Query Match	24.8%;	Score 81;	DB 10;	Length 1125;
Best Local Similarity	46.08;	Pred. No. 0.14;		
Matches 144;	Conservative 28;	Mismatches 141;	Indels 0;	Gaps 0

Qy 15 aaatttagcacagaagaatgltttaacgcgaataaagataatacaaaagatatattaattat 74
||| | | | | | : | : | : | | | | | | : :
Db 1079 AAAAAAAAAAAAAAATTTTTTTTWTWWTTTTWNNAAAAAAAAAAAAAAAAAAAWAAMW 1020

Oy 75 atgctgaataattacatcatatgatgaagtatagttcaataataaaagtgttcctcggg 134
 | || :||: || | :| | | | | :| | |
Db 1019 TAAAAAAATATWWTWTTTATTATTAATAAAAAAAAAAATAATATATWAAAAAATAWA 960

```

Oy      135  ggaacacttttttggttttaaaaaaggaatatataataaatctagatataaagtcgtaaaata 194
        ||::|||::||| ||| ||| ||| |||:::| ||| |||::|
Db      959  wwtattttttttttttttatataaaaaaaawwawwawwatttttttaaaaaaaawwaa 900

```

Qy 195 atcatttcattcaaatcgtttaaaaattgcatacaattgattgtaaaaaaaattcca 254
| | | | | : | | | | | : | | | | | :
Db 899 AAAAAATAAATTTTNTAATAAAMWTAATAAAAAATANTTTATWAAAAAATAAAW 840

Oy 255 ggggggaataataatgaaaaaaatatattccaagttactgtcaattttaagtctcatg 314
|||::||: ||: | ||: || | || | ||| | |||
Db 839 AAAAAAAATTTAATAAAAANATTAATTAWAAAAAATTTTTNANTATTTTTTTTTTNN 780

Qy	315	tttcttatgtt	327
Db	779	TATTTTNNTTTT	767

RESULT 14
CNS020K7

DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone 222L11 of library G from Tetraodon nigroviridis, genomic survey sequence.

VERSION ALL175696.1 GI:7813753
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis.

Eukaryota; Metazoa; Craniata, Vertebrata; Euteleostomi
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;

REFERENCE
AUTHORS
1 (bases 1 to 1092)
Roest-Crollius, H., Jallion, O., Dasilva, C., Fizames, C., Fisher, C.
Bonneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and
Bouneau, L.

TITLE	JOURNAL	DATE
Characterization and repeat analysis of the compact genome of the freshwater pufferfish <i>Tetraodon nigroviridis</i>	Unpublished	2 (bases 1 to 1000)

AUTHORS
Roest-Crollius, H., Jallion, O., Dasilva, C., Bouneau, L., Fisher, C.,
Bernot, A., Fizames, C., Wincker, P., Brotier, P., Quetier, F.,
Saurin, W. and Weissenbach, J.

Tetraodon nigroviridis DNA sequence
Unpublished
3 (bases 1 to 1092)

TITLE	Direct Submission
JOURNAL	Submitted (12-APR-2000) to the EMBL/Genbank/DBJ databases
COMMENT	This sequence is a single read and was generated as part of a 1

genome. For more information, please take a look at <http://www.genoscope.cns.fr/Tretraodon>.

Location/Qualifiers

```

/organism="Neitradon nigroviridis"
/db_xref="taxon:99883"
/clone="222111"
/clone_11b="G"
/notes="densoseq sequence ID : CONG222CF06LPI-end : T7
BASE COUNT      383 a      169 c      165 g      262 t      113 others
ORIGIN

```

Query Match	24.6%;	Score 80.6;	DB 13;	Length 1092;
Best Local Similarity	41.2%;	Pred. No. 0.16;		
Matches 132;	Conservative 60;	Mismatches 125;	Indels 3;	Gaps 1

```

Qy      8 acatcttaaattagcgcagaaagtgtttaaayaaataagatataaaaaagatatat 67
        ||:|:|:::| | | | | | | | | | | | | | | | | | | | | |
Db 619 ATWTTTAAWWWWAAAAAANNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 678

```

```

Qy 68 taattatagctgaaaaatttataatgataagtaagtttaataataaaaaagct 12
    ||| ||| ||| ||| ||| : || : ||| :
Db 679 AAAAAAAAAATTTTTTTTTTTTTTAAAAAAAAAAAAAAAAAAATTAATAAATAA 738

```

```
QY      128 tctcgggagacacttttltgtttaaaaaagaataataaattcagataaaatg 18
          : | ||:|| |::: ||| || :::|| | |:
Db      739 AAWAAWAAWWAAWTWTTTTAAWWWWWAAAA--NAAAWWWTATATTTTTAAMWA 79
```

Qy 188 taaataatctatttctatttaaaccttcgttaaanaattgataaatggaatcgttaaaaa 24
|:::| | | : |::| : | ::|| | | |
Db 796 TTAATTTTAAATAAANAATTAAATTAATAAAAAAATAATTAATTTTAAAN 85

Oy 248 aatttcagggggaatacataagyaaaaaaaatlattlccaagtltactcgtaatlttcaagc 30
|::| : : | | | | | | | | : : : : | | : : |
Db 856 AWTTTAAWAAATTTWWAAGTWTAAAAAAAATTAAAAAAWAWMTTTTTTTTTTWTAAMWT 913

```

QY      308  ttcatgtttctctatcgtt 327
          :| | | | | | | | | |
Db      916  WTAAATWTTTTTTTTTTTTT 935

```

RESULT 15
CNS0080F/C

DEFINITION
Drosophila melanogaster genome survey sequence TERT end of BACR17022 of RPCI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.

```
VERSION      AL052232.1  GI:4933983
KEYWORDS
SOURCE      GSS.
            fruit fly.
            fly.
```

Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (02-JUN-1999) Genoscope - Centre National de Sequençage
RD 101 01006 Evry - Evry - FRANCE (E-mail : Genoscope@genoscope.cnr.fr)

COMMENT

- Web : www.genoscope.cns.fr

Determination of this BAC-end sequence was carried out as part of the BAC-end sequencing project of the Berkeley Drosophila Genome Project (BDGP) in collaboration with the Berkeley Drosophila Genome Project (BDGP).

melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP *Drosophila* melanogaster BAC library was prepared by Kazuo Ito. Osseogawa and

Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of *Drosophila* DNA provided by the BDGP from the

P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, and filters for hybridization from the BACPAC Resource Center can be found at <http://bacpac.med.utoronto.ca/bacpac>.

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:50:52 ; Search time 216.42 Seconds
(Without alignments)
1295.376 Million cell updates/sec

Title: US-09-531-438-3
Perfect score: 327
Sequence: 1 atttgagatcttaattt.....tttcattgttctattgtt 327

Scoring table: IDENTITY_NIC
Gapop 10.0 , Gapept 1.0

Searched: 930621 seqs, 428662619 residues
Total number of hits satisfying chosen parameters: 1861242

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: N_Geneseq_1101:*
2: /SIDS2/gcgdata/geneseq/geneseq/NA1980.DAT:*
3: /SIDS2/gcgdata/geneseq/geneseq/NA1981.DAT:*
4: /SIDS2/gcgdata/geneseq/geneseq/NA1982.DAT:*
5: /SIDS2/gcgdata/geneseq/geneseq/NA1983.DAT:*
6: /SIDS2/gcgdata/geneseq/geneseq/NA1984.DAT:*
7: /SIDS2/gcgdata/geneseq/geneseq/NA1985.DAT:*
8: /SIDS2/gcgdata/geneseq/geneseq/NA1986.DAT:*
9: /SIDS2/gcgdata/geneseq/geneseq/NA1987.DAT:*
10: /SIDS2/gcgdata/geneseq/geneseq/NA1988.DAT:*
11: /SIDS2/gcgdata/geneseq/geneseq/NA1989.DAT:*
12: /SIDS2/gcgdata/geneseq/geneseq/NA1990.DAT:*
13: /SIDS2/gcgdata/geneseq/geneseq/NA1991.DAT:*
14: /SIDS2/gcgdata/geneseq/geneseq/NA1992.DAT:*
15: /SIDS2/gcgdata/geneseq/geneseq/NA1993.DAT:*
16: /SIDS2/gcgdata/geneseq/geneseq/NA1994.DAT:*
17: /SIDS2/gcgdata/geneseq/geneseq/NA1995.DAT:*
18: /SIDS2/gcgdata/geneseq/geneseq/NA1996.DAT:*
19: /SIDS2/gcgdata/geneseq/geneseq/NA1997.DAT:*
20: /SIDS2/gcgdata/geneseq/geneseq/NA1998.DAT:*
21: /SIDS2/gcgdata/geneseq/geneseq/NA1999.DAT:*
22: /SIDS2/gcgdata/geneseq/geneseq/NA2000.DAT:*
23: /SIDS2/gcgdata/geneseq/geneseq/NA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	327	100.0	327	20	AAK60300
2	327	100.0	1392	20	AAK60299
3	130.2	39.8	936	22	AAF58252
4	130.2	39.8	936	22	AAF58254
5	130.2	39.8	936	22	AAF58257
6	130.2	39.8	936	22	AAF58259
7	130.2	39.8	936	22	AAF58262
8	130.2	39.8	936	22	AAF58255
9	129.8	39.7	936	22	AAF58252
10	129.8	39.7	936	22	AAF58254
11	129.8	39.7	936	22	AAF58257

C 12	129.8	39.7	936	22	AAF58259
C 13	129.8	39.7	936	22	AAF58262
C 14	129.8	39.7	936	22	AAF58255
C 15	91.6	28.0	244	22	AAF58238
C 16	90.8	27.8	244	22	AAF58238
C 17	66.8	20.4	4590	22	AAF58238
C 18	66.4	20.3	2486	21	AAAB0472
C 19	64.8	19.8	20674	21	AAAB07037
C 20	63.8	19.5	2435	21	AAAC58017
C 21	63.2	19.3	9789	17	AAAT3865
C 22	62.2	19.0	6243	20	AAAT4852
C 23	61.6	18.8	19124	18	AAAT09598
C 24	61.6	18.8	19124	21	AAAT72882
C 25	61	18.7	700	22	AAAT98287
C 26	61	18.7	5852	12	AAAT93026
C 27	60.4	18.5	700	22	AAAT11710
C 28	60.2	18.4	1341	20	AAAT3026
C 29	60.2	18.4	1410	20	AAAT1756
C 30	59.6	18.2	513445	22	AAAT1755
C 31	59.4	18.2	6265	20	AAAT1373
C 32	59	18.0	381	22	AAAT08523
C 33	58.8	18.0	53585	20	AAAT93355
C 34	58.4	17.9	3975	9	AAAT0251
C 35	58.4	17.9	3975	13	AAAT1157
C 36	58.4	17.9	163319	21	AAAT22999
C 37	58.2	17.8	1132	21	AAAT2306
C 38	58.2	17.8	2503	15	AAAT12929
C 39	58.2	17.8	9048	18	AAAT5480
C 40	57.8	17.7	8310	20	AAAT43225
C 41	57.6	17.6	1907	20	AAAT29911
C 42	57.4	17.6	366	22	AAAT20307
C 43	57	17.4	20674	21	AAAT3356
C 44	56.8	17.4	605	17	AAAT58017
C 45	56.8	17.4	665	21	AAAT31530
C 45	56.8	17.4	665	21	AAAT31996

ALIGNMENTS

RESULT 1	
ID AAK60300	standard; DNA; 327 BP.
XX AC AAK60300:	
XX AC 12-AUG-1999	(first entry)
XX DT	
XX DE	Promoter of the beta-2 toxin gene of Clostridium perfringens type C.
XX DE	
XX KW	Beta-2 toxin; Clostridium perfringens type C; gene promoter;
XX KW	vaccine; Clostridium tetani; ss.
XX OS	Clostridium perfringens.
XX OS	
XX PN	FR2768747-A1.
XX PD	26-MAR-1999.
XX PF	19-SEP-1997; 97FR-0011710.
XX PF	
XX PR	19-SEP-1997; 97FR-0011710.
XX PA	(INSP) INST PASTEUR.
XX PA	
XX PL	Gilbert M, Popoff MR;
XX DR	WPI; 1999-217498/19.
XX PT	Clostridium beta2 toxin gene promoter and signal sequence - useful
XX PT	against toxins from Clostridium perfringens
XX PS	Claim 1; Page 32; 46pp; French.
XX PS	

Oligonucleotide D2
Oligonucleotide D2
Oligonucleotide D1
Oligonucleotide D1
Oligonucleotide D1
Sequence encoding
Nucleotide sequenc
Arachidonic acid m
Beta glucosidase (
CDNA encoding Plas
Clostridium specie
Plasmodium var-7 g
Plasmodium var-7 p
Human inflammatory
Dictyostellium plas
Human inflammatory
B. burgdorferi ant
B. burgdorferi ant
Soybean 318013 reg
NBP46 (root lectin
Human chromosome 1
Borrelia burgdorfe
Malaria-specific g
SERP gene. Plasm
Arabidopsis thalia
Aspergillus oryzae
pNPX30 xylanase cd
Brassica napus FCA
CDNA encoding a SC
Borrelia burgdorfe
Human chromosome 1
Arachidonic acid m
Human 3' apolipop
Human apolipoprote

CC The present sequence represents the promoter of the beta-2 toxin
CC gene of Clostridium perfringens type C. The beta2-toxin promoter
CC and gene sequences can be used to produce vaccines against Clostridium,
CC and especially Clostridium perfringens, or Clostridium
CC tetani.

CC Sequence 327 BP; 141 A; 13 C; 44 G; 129 T; 0 other;

Query Match 100.0%; Score 327; DB 20; Length 327;

Best Local Similarity 100.0%; Pred. NO. 2.2e-35;
Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```
QY 1 attgggatatcttaatttagcacagaagaatgtttaataaagaataaaaaa 60
DB 1 attgggatatcttaatttagcacagaagaatgtttaataaagaataaaaa 60
QY 61 gataataataatagctggaatttaataatagatagtaataataa 120
DB 61 gataataataatagctggaatttaataatagatagtaataataa 120
QY 121 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 180
DB 121 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 180
QY 181 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 240
DB 181 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 240
QY 241 taaaaaaatttcagggggaataataatgaaaaaattattccaagtctgaatt 300
DB 241 taaaaaaatttcagggggaataataatgaaaaaattattccaagtctgaatt 300
QY 301 ttatggtttcatggttttcttattgt 327
DB 301 ttatggtttcatggttttcttattgt 327
```

RESULT 2

AAK60299 standard; DNA; 1392 BP.

AAK60299;

12-AUG-1999 (first entry)

DNA encoding the beta-2 toxin of Clostridium perfringens type C.

Beta-2 toxin; Clostridium perfringens type C; gene promoter;
KW vaccine; Clostridium tetani; ss.

Clostridium perfringens.

FR2768747-A1.

26-MAR-1999.

19-SEP-1997; 97FR-0011710.

19-SEP-1997; 97FR-0011710.

(INSP) INST PASTEUR.

Gibert M, Popoff MR.

WPI, 1999-217498/19.

P-PSDB; AAY16591.

Clostridium beta2 toxin gene promoter and signal sequence - useful
PT against toxins from Clostridium perfringens

Example A; Page 31; 46pp; French.

CC The present sequence encodes the beta-2 toxin of Clostridium
CC perfringens type C. The specification describes the Clostridium
CC perfringens beta 2 toxin gene promoter (see AAK60300). The
CC sequences can be used to produce vaccines against Clostridium,
CC and especially Clostridium perfringens, or Clostridium
CC tetani.

CC Sequence 1392 BP; 606 A; 115 C; 209 G; 462 T; 0 other;

Query Match 100.0%; Score 327; DB 20; Length 1392;

Best Local Similarity 100.0%; Pred. NO. 1.7e-35;
Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```
QY 1 attgggatatcttaatttagcacagaagaatgtttaataaagaataaaaaa 60
DB 1 attgggatatcttaatttagcacagaagaatgtttaataaagaataaaaaa 60
QY 61 gataataataatagctggaatttaataatagatagtaataataa 120
DB 61 gataataataatagctggaatttaataatagatagtaataataa 120
QY 121 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 180
DB 121 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 180
QY 181 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 240
DB 181 aaagtgcttcgaggacaccttttggtttaaaaggaataataaataattagat 240
QY 241 taaaaaaatttcagggggaataataatgaaaaaattattccaagtctgaatt 300
DB 241 taaaaaaatttcagggggaataataatgaaaaaattattccaagtctgaatt 300
QY 301 ttatggtttcatggttttcttattgt 327
DB 301 ttatggtttcatggttttcttattgt 327
```

RESULT 3

AAF58252 standard; DNA; 936 BP.

AAF58252;

24-APR-2001 (first entry)

Oligonucleotide D1835.

Electron-transfer group; ETM; mismatch; genotyping;
KW gene expression; ss.

Synthetic.

WO200107665-A2.

01-FEB-2001.

26-JUL-2000; 2000WO-US20476.

26-JUL-1999; 99US-0145695.

17-MAR-2000; 2000US-0190259.

(CLIN-) CLINICAL MICRO SENSORS INC.

Umek RM;

WPI, 2001-159728/16.

Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
a single surface

PI Umek RM;
 XX WPI: 2001-159728/16.
 XX Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface -
 XX
 XX Example 6: Page 128; 159pp: English.
 XX
 XX The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 CC
 XX Sequence 936 BP; 5 A; 139 C; 10 G; 6 T; 776 other;

Query Match 39.8%; Score 130.2; DB 22; Length 936;
 Best Local Similarity 0.9%; Pred. No. 9.2e-10;
 Matches 3; Conservative 268; Mismatches 56; Indels 0; Gaps 0;
 QY 1 attgggataccttaattagcacagaagaatgtttaaatgaataaagataaataa 60
 DB 136 www..... 195
 QY 61 gatataatataatagctgaataattataatataatgataagataatgaataa 120
 DB 196 gwww..... 255
 QY 121 aaaggtctcgggggacacttttgtttaaaaggaataataaattagat 180
 DB 256 www..... 315
 QY 181 aaagtgtaataataatttttaatttaattgttaaaattgataataattgattg 240
 DB 316 www..... 375
 QY 241 taaaaaaatttcagggggaataataaataaattattcaagttcactgtaatt 300
 DB 376 www..... 435
 QY 301 ttatgtttcatgtttctctatggt 327
 DB 436 www..... 462

RESULT 8
 AAF58255
 ID AAF58255 standard; DNA; 938 BP.
 XX
 AC AAF58255;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D1876.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 XX
 OS Synthetic.
 XX
 PN WO200107665-A2.
 XX
 PD 01-FEB-2001.
 XX
 PD 26-JUL-2000; 2000WO-US20476.
 PF
 XX 26-JUL-1999; 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX

PA (CLIN-) CLINICAL MICRO SENSORS INC.
 XX
 XX Umek RM;
 XX WPI: 2001-159728/16.
 XX Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface -
 XX
 XX Example 6: Page 127; 159pp: English.
 XX
 XX The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 CC
 XX Sequence 938 BP; 4 A; 144 C; 9 G; 5 T; 776 other;

Query Match 39.8%; Score 130.2; DB 22; Length 938;
 Best Local Similarity 0.9%; Pred. No. 9.2e-10;
 Matches 3; Conservative 268; Mismatches 56; Indels 0; Gaps 0;
 QY 1 attgggataccttaattagcacagaagaatgtttaaatgaataaagataaataa 60
 DB 136 www..... 195
 QY 61 gatataatataatagctgaataattataatataatgataagataatgaataa 120
 DB 196 gwww..... 255
 QY 121 aaaggtctcgggggacacttttgtttaaaaggaataataaattagat 180
 DB 256 www..... 315
 QY 181 aaagtgtaataataatttttaatttaattgttaaaattgataataattgattg 240
 DB 316 www..... 375
 QY 241 taaaaaaatttcagggggaataataaataaattattcaagttcactgtaatt 300
 DB 376 www..... 435
 QY 301 ttatgtttcatgtttctctatggt 327
 DB 436 www..... 462

RESULT 9
 AAF58252/c
 ID AAF58252 standard; DNA; 936 BP.
 XX
 AC AAF58252;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D1835.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 XX
 OS Synthetic.
 XX
 PN WO200107665-A2.
 XX
 PD 01-FEB-2001.
 XX
 PD 26-JUL-2000; 2000WO-US20476.
 PF
 XX 26-JUL-1999; 99US-0145695.
 PR

PR 17-MAR-2000: 2000US-0190259.
XX
PA (CLIN-) CLINICAL MICRO SENSORS INC.
XX
PI
XX
XX
DR WPI: 2001-159728/16.
XX
XX
PT Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
PT a single surface -
XX
XX
PS Example 6; Page 127; 159pp; English.
XX
XX The present invention relates to a composition comprising two nucleic
CC acids each containing an electron-transfer group (ETM) having
CC different redox potentials. The invention is used for electronic
CC detection of nucleic acids, especially of substitutions (mismatches)
CC and single-nucleotide polymorphisms, e.g. for genotyping,
CC monitoring gene expression.
XX
XX Sequence 936 BP; 4 A; 139 C; 10 G; 7 T; 776 other;
XQ

XX 26-JUL-1999: 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX
 XX (CLIN-) CLINICAL MICRO SENSORS INC.
 PA
 XX
 XX Umek RM;
 PI
 XX
 DR WPI; 2001-159728/16.
 XX
 PT Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface
 XX
 XX
 PS Example 6; Page 127; 159pp; English.
 XX
 XX The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 XX
 XX Sequence 936 BP; 4 A; 144 C; 7 G; 5 T; 776 other;

XX 26-JUL-2000; 2000MO-US20476.
 PF 26-JUL-1999; 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX
 PA (CLIN-) CLINICAL MICRO SENSORS INC.
 XX
 PI Umek RM;
 DR WPI; 2001-159728/16.
 XX
 PT Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface
 XX
 PS Example 6; Page 127; 159pp; English.
 XX
 CC The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 XX
 SO Sequence 936 BP; 5 A; 142 C; 7 G; 6 T; 776 other;

Query Match 39.7%; Score 129.8; DB 22; Length 936;
 Best Local Similarity 0.6%; Pred. No. 1e-09;
 Matches 2; Conservative 269; Mismatches 56; Indels 0; Gaps 0;
 OY 1 attgggatacttaatttagcacagaagaatgtttaatgaataaagaataaataa 60
 DB 750 ww 691
 OY 61 gataataataatagctgaaattataataatagataagtaataataa 120
 DB 690 Gww 631
 OY 121 aaagtctctcgaggacactttttgtttaaaagaataataaattagat 180
 DB 630 Www 571
 OY 181 aaagtgtaaataatttttatttaattgtaaaattgataataattgattg 240
 DB 570 Www 511
 OY 241 taaaaaaattcagggggaataataatgaataaattattcaagtttaactt 300
 DB 510 Www 451
 OY 301 ttatgtttcattgtttcttattgtt 327
 DB 450 Wwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwww 424

RESULT 12
 AAF58259/C
 ID AAF58259 standard; DNA; 936 BP.
 AC AAF58259;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D2004.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 OS Synthetic.
 XX
 PN WO200107665-A2.

XX 01-FEB-2001.
 PD 26-JUL-2000; 2000MO-US20476.
 XX
 PF 26-JUL-1999; 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX
 PA (CLIN-) CLINICAL MICRO SENSORS INC.
 XX
 PI Umek RM;
 DR WPI; 2001-159728/16.
 XX
 PT Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface
 XX
 PS Example 6; Page 128; 159pp; English.
 XX
 CC The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 XX
 SO Sequence 936 BP; 6 A; 138 C; 8 G; 8 T; 776 other;

Query Match 39.7%; Score 129.8; DB 22; Length 936;
 Best Local Similarity 0.6%; Pred. No. 1e-09;
 Matches 2; Conservative 269; Mismatches 56; Indels 0; Gaps 0;
 OY 1 attgggatacttaatttagcacagaagaatgtttaatgaataaagaataaataa 60
 DB 750 ww 691
 OY 61 gataataataatagctgaaattataataatagataagtaataataa 120
 DB 690 Gww 631
 OY 121 aaagtctctcgaggacactttttgtttaaaagaataataaattagat 180
 DB 630 Www 571
 OY 181 aaagtgtaaataatttttatttaattgtaaaattgataataattgattg 240
 DB 570 Www 511
 OY 241 taaaaaaattcagggggaataataatgaataaattattcaagtttaactt 300
 DB 510 Www 451
 OY 301 ttatgtttcattgtttcttattgtt 327
 DB 450 Wwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwww 424

RESULT 13
 AAF58262/C
 ID AAF58262 standard; DNA; 936 BP.
 AC AAF58262;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D2007.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 OS Synthetic.

```

XX WO200107665-A2.
XX
XX 01-FEB-2001.
XX
XX 26-JUL-2000; 2000WO-US20476.
XX
XX 26-JUL-1999; 99US-0145695.
XX PR 17-MAR-2000; 2000US-0190259.
XX
XX (CLIN-) CLINICAL MICRO SENSORS INC.
XX
XX Umek RM:
XX
XX WPI; 2001-159728/16.
XX
XX Nucleic acids containing electron-transfer group, useful as labels in
XX hybridization assays, e.g. for genotyping, allowing repeat analyses on
XX a single surface.
XX
XX Example 6; Page 128; 159pp; English.
XX
XX The present invention relates to a composition comprising two nucleic
XX acids each containing an electron-transfer group (ETM) having
XX different redox potentials. The invention is used for electronic
XX detection of nucleic acids, especially of substitutions (mismatches)
XX and single-nucleotide polymorphisms, e.g. for genotyping,
XX monitoring gene expression.
XX
XX Sequence 936 BP; 5 A; 139 C; 10 G; 6 T; 776 other;
XX

```

```

Query Match          39.7%; Score 129.8; DB 22; Length 936;
Best Local Similarity 0.6%; Pred. No. 1e-09; Mismatches 56; Indels 0; Gaps 0;
Matches 2; Conservative 269;

QY 1 attgggatacttaaatagcacagaagaatgtttaaagaataaataaataa 60
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 750 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 691
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 61 gatatacttaataagctgaataattataatgataagtagttaataata 120
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
DB 690 GWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 631
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 121 aaagcttcgaggacacttttctttaaaagaaataataaataatagat 180
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 630 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 571
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 181 aaagctgtaaaataatttataattgtaaaattgataataattgaatg 240
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 570 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 511
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 241 taaaaaaatttcaggagggaataataatgaaaaaaattattccaagt 300
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 510 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 451
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 301 ttatgtttcattgtttcttattgt 327
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 450 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 424
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|

```

```

RESULT 14
AAF58255/C
ID AAF58255 standard; DNA; 938 BP.
XX
XX AAF58255;
XX
XX 24-APR-2001 (first entry)
XX
XX Oligonucleotide D1876.
XX
XX Electron-transfer group; ETM; mismatch; genotyping;
XX gene expression; ss.
XX

```

```

XX OS Synthetic.
XX
XX WO200107665-A2.
XX
XX 01-FEB-2001.
XX
XX 26-JUL-2000; 2000WO-US20476.
XX
XX 26-JUL-1999; 99US-0145695.
XX PR 17-MAR-2000; 2000US-0190259.
XX
XX (CLIN-) CLINICAL MICRO SENSORS INC.
XX
XX Umek RM:
XX
XX WPI; 2001-159728/16.
XX
XX Nucleic acids containing electron-transfer group, useful as labels in
XX hybridization assays, e.g. for genotyping, allowing repeat analyses on
XX a single surface.
XX
XX Example 6; Page 127; 159pp; English.
XX
XX The present invention relates to a composition comprising two nucleic
XX acids each containing an electron-transfer group (ETM) having
XX different redox potentials. The invention is used for electronic
XX detection of nucleic acids, especially of substitutions (mismatches)
XX and single-nucleotide polymorphisms, e.g. for genotyping,
XX monitoring gene expression.
XX
XX Sequence 938 BP; 4 A; 144 C; 9 G; 5 T; 776 other;
XX

```

```

Query Match          39.7%; Score 129.8; DB 22; Length 938;
Best Local Similarity 0.6%; Pred. No. 1e-09; Mismatches 56; Indels 0; Gaps 0;
Matches 2; Conservative 269;

QY 1 attgggatacttaaatagcacagaagaatgtttaaagaataaataaataa 60
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 750 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 691
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 61 gatatacttaataagctgaataattataatgataagtagttaataataa 120
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
DB 690 GWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 631
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 121 aaagcttcgaggacacttttctttaaaagaaataataaataatagat 180
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 630 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 571
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 181 aaagctgtaaaataatttataattgtaaaattgataataattgaatg 240
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 570 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 511
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 241 taaaaaaatttcaggagggaataataatgaaaaaaattattccaagt 300
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 510 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 451
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
QY 301 ttatgtttcattgtttcttattgt 327
   :::: :::: :::: :::: :::: :::: :::: :::: :::: ::::
DB 450 WWWWWWMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMMM 424
   |:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|

```

```

RESULT 15
AAF58238
ID AAF58238 standard; DNA; 244 BP.
XX
XX AAF58238;
XX
XX 24-APR-2001 (first entry)
XX
XX Oligonucleotide D1250; D1102.
XX

```

KW Electron-transfer group; ETM; mismatch; genotyping;
KW gene expression; ss.
VV

gene expression; ss.

Synthetic.

PN W0200107665-A2.

01-FEB-2001.

PF 26-JUL-2000; 2000WO-US20476.

PR 26-JUL-1999; 99US-0145695.

[illegible]

XX

XX
XX
FIFTY-FOUR
2000

XX 33

PT Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
PT a single surface -

PS Example 4; Page 120; 159pp; English.

CC The present invention relates to a

CC different redox potentials. The inve

CC and single-nucleotide polymorphisms, e.g. for genotyping,

XX

Query Match	28.0%;	Score 91.6;	DB 22;	Length 244;
-------------	--------	-------------	--------	-------------

Matches 10; Conservative 167; Mismatches 31; Indels 0; Gaps 0;

QY 42 gaaataagataataaaaagatatattaattatagctgaaatttataattatgat 101

[illegible]

QY 102 aagtatagtttaataataaaagtgttctcgggggacacttttctgttttaaaaag

Db 67 www.wtwh.com C www.wtwh.com 126

QY 162 atataataaatttagataaaagtgtaaaataattatttttaaatltgttaaaa 221

[illegible]

QY 222 attgataataattgaattgtaaaaaa 249 .

Db 187 www.wtlttaagaca 214

Search completed: January 24, 2002, 02:22:17
Job #1985-1005

Search completed: January 24, 2002, 02:22:17
Job time: 1885 sec

Job time: 1885 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:51:38 ; Search time 2099.46 Seconds
(without alignments)
1673.702 Million cell updates/sec

Title: US-09-531-438-3
Perfect score: 327
Sequence: 1 atttggatattcttaattt.....tttcattttcttattgtt 327

Scoring table: OLIGO-MNC
Gapop 60.0, Gapext 60.0

Searched: 11351937 seqs, 5372889281 residues

Word size: 0

Total number of hits satisfying chosen parameters: 80718

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

EST:*
1: em_estfun:*
2: em_esthm:*
3: em_estin:*
4: em_estom:*
5: em_estpl:*
6: em_estba:*
7: em_estro:*
8: em_estov:*
9: em_hic:*
10: gb_estl:*
11: gb_est2:*
12: gb_hic:*
13: gb_gss:*
14: em_gss_fun:*
15: em_gss_hum:*
16: em_gss_inv:*
17: em_gss_pln:*
18: em_gss_pro:*
19: em_gss_rod:*
20: em_gss_vrt:*
21: em_gss_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	4.6	42	10	AU011968 AU011968
2	15	4.6	42	10	AU011969 AU011969
3	15	4.6	42	10	AU011971 AU011971
4	15	4.6	42	10	AU011973 AU011973
5	15	4.6	49	10	AA922891 015105.5
6	15	4.6	50	11	C01094 HUMGS000775
7	14	4.3	24	13	AZ781748 2M0021M18
8	14	4.3	28	11	C01204 HUMGS000790
9	14	4.3	34	10	AU038857 AU038857
10	14	4.3	42	10	AW333885 S27E6 AGS
11	14	4.3	42	13	AZ634761 IM0490C17
12	14	4.3	46	13	AZ459612 IM0264002

C 13	14	4.3	47	13	AZ345468
14	14	4.3	50	10	BE043289
15	14	4.3	50	13	TA154F10Q
C 16	13	4.0	19	13	AZ331628
C 17	13	4.0	25	13	AZ829725
C 18	13	4.0	26	13	AZ309204
C 19	13	4.0	26	13	AZ866662
C 20	13	4.0	26	13	TA123B12Q
C 21	13	4.0	27	13	AZ784620
C 22	13	4.0	28	13	AZ452653
C 23	13	4.0	30	13	AZ623794
C 24	13	4.0	32	11	H40874
C 25	13	4.0	32	13	AZ458690
C 26	13	4.0	34	10	AA906810
C 27	13	4.0	34	13	AZ586746
C 28	13	4.0	34	13	AZ781725
C 29	13	4.0	35	10	AW246486
C 30	13	4.0	35	11	D45807
C 31	13	4.0	36	13	AZ314238
C 32	13	4.0	37	10	AA913140
C 33	13	4.0	37	10	AU009123
C 34	13	4.0	40	10	AA916625
C 35	13	4.0	40	10	AA922076
C 36	13	4.0	41	11	C00434
C 37	13	4.0	41	13	AZ777050
C 38	13	4.0	43	13	AZ371136
C 39	13	4.0	43	13	AZ464392
C 40	13	4.0	43	13	AZ575514
C 41	13	4.0	45	11	D20668
C 42	13	4.0	46	13	AZ459612
C 43	13	4.0	46	13	AZ834972
C 44	13	4.0	46	13	AZ991460
C 45	13	4.0	49	10	AI630064

ALIGNMENTS

RESULT 1
AU011968
LOCUS AU011968 42 bp mRNA EST 03-AUG-1998
DEFINITION AU011968 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA clone spc06162, mRNA sequence.
AU011968
ACCESSION AU011968.1 GI:3356877

VERSION
KEYWORDS
SOURCE
ORGANISM
EST.
fission yeast.
Schizosaccharomyces pombe
Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetales; Schizosaccharomycetaceae;
Schizosaccharomyces.
1 (bases 1 to 42)

REFERENCE
AUTHORS
TITLE
Moriyomo,M. and Mita,K.
Identification of expressed sequence tags of Schizosaccharomyces
pombe
(1998)

JOURNAL
COMMENT
Unpublished
Contact: Mitsueki Moriyomo
Genome Research Group
National Institute of Radiological Sciences
9-1, Anagawa-4-chome, Inage-Ku, Chiba 263-8555, Japan
Email: moriyomo@nirs.go.jp

FEATURES

source

1..42
/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06162"
/sex="h minus"
/note="Vector: M13mp19. The cDNA library of
Schizosaccharomyces pombe was prepared by cloning cDNA
into the SmaI site of M13mp19 DNA and the direction of DNA
sequences was not always from 5' to 3'. The cDNA data of

Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)

Query Match 4.6%; Score 15; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 9.6e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 142 ttctgttttaaaaa 156
|||||
Db 22 TTTTGTGTTTAAAAA 36

RESULT 2

LOCUS AU011969 42 bp mRNA EST 03-AUG-1998
DEFINITION AU011969 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA clone spc06163, mRNA sequence.

ACCESSION AU011969
VERSION AU011969.1 GI:3356878

KEYWORDS EST.
SOURCE fission yeast.

ORGANISM Schizosaccharomyces pombe

Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetales; Schizosaccharomycetaceae;

Schizosaccharomyces.

1 (bases 1 to 42)

Moriyomo, M. and Mita, K.

Identification of expressed sequence tags of Schizosaccharomyces pombe

JOURNAL Unpublished (1998)
COMMENT Contact: Mitsuoki Moriyomo

Genome Research Group

National Institute of Radiological Sciences

9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan

Email: moriyomo@nirs.go.jp.

Location/Qualifiers

FEATURES

source 1.42

/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06163"

/sex="h minus"
/note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe late log phase cDNA"

into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)

BASE COUNT 19 a 1 c 7 g 15 t

ORIGIN

Query Match 4.6%; Score 15; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 9.6e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 142 ttctgttttaaaaa 156
|||||
Db 22 TTTTGTGTTTAAAAA 36

RESULT 3

LOCUS AU011971 42 bp mRNA EST 03-AUG-1998
DEFINITION AU011971 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA clone spc06165, mRNA sequence.

ACCESSION AU011971
VERSION AU011971.1 GI:3356880

KEYWORDS EST.

SOURCE

ORGANISM

fission yeast.

Schizosaccharomyces pombe

Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;

Schizosaccharomycetales; Schizosaccharomycetaceae;

Schizosaccharomyces.

1 (bases 1 to 42)

Moriyomo, M. and Mita, K.

Identification of expressed sequence tags of Schizosaccharomyces pombe

JOURNAL Unpublished (1998)

COMMENT Contact: Mitsuoki Moriyomo

Genome Research Group

National Institute of Radiological Sciences

9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan

Email: moriyomo@nirs.go.jp.

Location/Qualifiers

FEATURES

source 1.42

/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06165"

/sex="h minus"
/note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe late log phase cDNA"

into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)

BASE COUNT 19 a 1 c 7 g 15 t

ORIGIN

Query Match 4.6%; Score 15; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 9.6e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 142 ttctgttttaaaaa 156
|||||
Db 22 TTTTGTGTTTAAAAA 36

RESULT 4

LOCUS AU011973 42 bp mRNA EST 03-AUG-1998

DEFINITION AU011973 Schizosaccharomyces pombe late log phase cDNA

Schizosaccharomyces pombe cDNA clone spc06167, mRNA sequence.

ACCESSION AU011973

VERSION AU011973.1 GI:3356882

KEYWORDS EST.

SOURCE fission yeast.

ORGANISM Schizosaccharomyces pombe

Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;

Schizosaccharomycetales; Schizosaccharomycetaceae;

Schizosaccharomyces.

1 (bases 1 to 42)

Moriyomo, M. and Mita, K.

Identification of expressed sequence tags of Schizosaccharomyces pombe

JOURNAL Unpublished (1998)

COMMENT Contact: Mitsuoki Moriyomo

Genome Research Group

National Institute of Radiological Sciences

9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan

Email: moriyomo@nirs.go.jp.

Location/Qualifiers

FEATURES

source 1.42

/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06167"

/sex="h minus"
/note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe late log phase cDNA"

into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)

BASE COUNT 19 a 1 c 7 g 15 t

ORIGIN

Query Match 4.6%; Score 15; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 9.6e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 142 ttctgttttaaaaa 156
|||||
Db 22 TTTTGTGTTTAAAAA 36

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 1000 Std Error: 0.00
 Plate: 0021 row: M column: 18
 Seq primer: CGTGTAAACGACGCGCAGT
 Class: Plasmid ends
 High quality sequence stop: 24.

FEATURES

source

1. 24
 Location/Qualifiers
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="U0022M0021M18"
 /clone_lib="Mouse 10Kb plasmid U0021M18"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g147321419b1AF129072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT

9 a 0 c 3 g 12 t

ORIGIN

Query Match 4.3%; Score 14; DB 13; Length 24;
 Best Local Similarity 100.0%; Pred. No. 2.9e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 198 attttattttaa 211

Db 7 ATTATTATTTTAA 20

RESULT 8

LOCUS

C01204 28 bp mRNA EST 23-JUL-1996

DEFINITION HUMS0007904 Human adult (K.Okubo) Homo sapiens cDNA, mRNA

ACCESSION C01204

VERSION C01204.1 GI:1433434

KEYWORDS EST.

SOURCE human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE

AUTHORS

Okubo, K.

BodyMap: human gene expression database

Unpublished (1995)

Contact: Okubo, K.

Institute for Molecular and Cellular Biol

1-3, Yamada-oka, Suita, Osaka Pref. 565, Japan

Tel: 06-877-5111(ex. 3315)

Email: kousaku@imcb.osaka-u.ac.jp

Human Gene Signature, 3'-directed cDNA sequence. We are not submitting the same cDNA sequence redundantly to DBJ since 1993. For the abundance information of clones with this sequence in this library and as well as in other 3'-directed libraries, see http://www.imcb.osaka-u.ac.jp/bodymap. The sequences of the clones represented by this GS sequences is also found there.

FEATURES

source

1. 28
 Location/Qualifiers
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="Human adult (K.Okubo)"
 /dev_stage="adult"
 /dev_stage="adult"

BASE COUNT

11 a 3 c 3 g 10 t 1 others

ORIGIN

Query Match 4.3%; Score 14; DB 11; Length 28;
 Best Local Similarity 100.0%; Pred. No. 2.7e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 198 attttattttaa 211

Db 20 ATTATTATTTTAA 7

RESULT 9

LOCUS

A0038857 34 bp mRNA EST 29-MAR-1999

DEFINITION A0038857 Dictyostelium discoideum SS (H.Urushihara) Dictyostelium

ACCESSION

A0038857

VERSION A0038857.1 GI:3985610

KEYWORDS EST.

SOURCE Dictyostelium discoideum.

ORGANISM

Dictyostelium discoideum.

REFERENCE

AUTHORS

Mori, T., Urushihara, H., Salto, T., Ugawa, Y., Mizuno, H., Yoshida, M.,

Yoshino, R., Mitra, B.N., Pl.M., Sato, T., Takemoto, K., Yasukawa, H.,

Williams, J., Maeda, M., Takeuchi, T., Ochiai, H. and Tanaka, Y.

The Dictyostelium developmental cDNA project: generation and

analysis of expressed sequence tags from the first-finger stage of

development

DNA Res. 5 (6), 335-340 (1998)

Contact: Hideko Urushihara

Institute of Biological Sciences

University of Tsukuba

3-3-10 Ten-nodai, Tsukuba, Ibaraki 305, Japan

Email: d402huesakura.cc.tsukuba.ac.jp

PROJECT = 'Dictyostelium discoideum cDNA project in Japan'

Location/Qualifiers

1. 34

/organism="Dictyostelium discoideum"

/strain="Ax4"

/db_xref="taxon:44689"

/clone="SSL566"

/clone_lib="Dictyostelium discoideum SS (H.Urushihara)"

/dev_stage="slug"

BASE COUNT 23 a 0 c 0 g 11 t

ORIGIN

Query Match 4.3%; Score 14; DB 10; Length 34;

Best Local Similarity 100.0%; Pred. No. 2.5e+05;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 110 ttaataataaaaa 123

Db 2 TTAATAATAAANA 15

RESULT 10


```

AW333885      42 bp      mRNA      EST      31-JAN-2000
LOCUS      AM333885
DEFINITION      S27B6 AGS-1 Pneumocystis carinii f. sp. carinii cDNA 3', mRNA
SEQUENCE
ACCESSION      AM333885
VERSION      AM333885.1 GI:6830242
KEYWORDS      EST.
SOURCE      Pneumocystis carinii f. sp. carinii.
ORGANISM      Eukaryota; Fungi; Ascomycota; Pneumocystidomycetes;
Pneumocystidaceae; Pneumocystis.
REFERENCE      1 (bases 1 to 42)
AUTHORS      Smilian,A.G., Arnold,J., Weise,M., Wunderlich,J., Staben,C., Edman
J.C., Kovacs,J. and Cushion,M.
JOURNAL      Expressed sequence tags from Pneumocystis carinii
COMMENT      Unpublished (2000)
CONTACT      Staben C
School of Biological Sciences
University of Kentucky
101 Morgan Building, University of Kentucky, Lexington, KY
40506-0225, USA
Tel: 606 257 2161
Fax: 606 257 1717
Email: staben@pop.uky.edu.
Location/Qualifiers
FEATURES
source
1..42
/organism="Pneumocystis carinii f. sp. carinii"
/db_xref="taxon:38081"
/clone_lib="AGS-1"
/lab_host="E. coli"
/note="Vector: lambda ZAP II; Site 1: EcoRI; Site 2: XhoI;
P. carinii organisms (3x10e9) from a single rat (99-1-6,
sacrificed on 3/17/99) at Cincinnati VA facilities.
Triool extracted RNA. Oligo dT priming, standard
conditions described by vendor, Stratagene. Further
details see www.uky.edu/Project/Pneumocystis/"
BASE COUNT      19 a      0 c      6 g      17 t
ORIGIN
Query Match      4.3%; Score 14; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 2.4e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY      164 ataataaattta 177
Db      3 ATTAATAAATTTA 16
RESULT 11
AZ634761/c      42 bp      DNA      GSS      13-DEC-2000
LOCUS      IM0490C17R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION      clone UUGC1M0490C17 R, DNA sequence.
ACCESSION      AZ634761
VERSION      AZ634761.1 GI:11756951
KEYWORDS      GSS.
SOURCE      house mouse.
ORGANISM      Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 42)
REFERENCE      1 (bases 1 to 42)
AUTHORS      Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.
JOURNAL      Mouse whole genome scaffolding with paired end reads from 10kb
COMMENT      Unpublished (2000)
CONTACT      Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

```

```

84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0490 row: C column: 17
Seq primer: CACACGAGGAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 42.
Location/Qualifiers
FEATURES
source
1..42
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone_lib="UUGC1M0490C17"
/lab_host="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/nares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD2 (g147321419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
BASE COUNT      22 a      5 c      4 g      11 t
ORIGIN
Query Match      4.3%; Score 14; DB 13; Length 42;
Best Local Similarity 100.0%; Pred. No. 2.4e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY      141 tttttgttttaa 154
Db      34 TTTTGTGTTTAA 21
RESULT 12
AZ459612/c      46 bp      DNA      GSS      04-OCT-2000
LOCUS      IM0264002R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION      clone UUGC1M0264002 R, DNA sequence.
ACCESSION      AZ459612
VERSION      AZ459612.1 GI:10617737
KEYWORDS      GSS.
SOURCE      house mouse.
ORGANISM      Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 46)
REFERENCE      1 (bases 1 to 46)
AUTHORS      Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.
JOURNAL      Mouse whole genome scaffolding with paired end reads from 10kb
COMMENT      Unpublished (2000)
CONTACT      Robert B. Weiss
University of Utah Genome Center

```

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0264 row: 0 column: 02
Seq primer: CACACAGAAACAGCATGACC
Class: plasmid ends
High quality sequence stop: 46.

FEATURES

source

1. 46
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C1M0264002"
/clone_1ib="Mouse 10kb plasmid U08C1M library"
/sex="Male"
/lab_host="E. coli strain XL10-gold, T1-resistant, F-"
/note="Vector: PMD42ny; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (g114732114|9b|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT
ORIGIN

21 a 5 c 4 g 16 t

Query Match 4.3%; Score 14; DB 13; Length 46;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 188 taaataattattt 201

Db 45 TAAATAATTATT 32

RESULT 13
AZ345468/C

LOCUS AZ345468 47 bp DNA GSS 29-SEP-2000
DEFINITION 1M0080N12F Mouse 10kb plasmid U08C1M library Mus musculus genomic
clone U08C1M080N12 F, DNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE

ORGANISM

house mouse.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE

1 (bases 1 to 47)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

TITLE

JOURNAL

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center

University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0080 row: N column: 12
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 47.

FEATURES

source

1. 47
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C1M080N12"
/clone_1ib="Mouse 10kb plasmid U08C1M library"
/sex="Male"
/lab_host="E. coli strain XL10-gold, T1-resistant, F-"
/note="Vector: PMD42ny; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (g114732114|9b|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT
ORIGIN

12 a 9 c 0 g 26 t

Query Match 4.3%; Score 14; DB 13; Length 47;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 112 aataataaaagt 125

Db 26 AATAATAAAAGT 13

RESULT 14
BE043289

LOCUS BE043289 50 bp mRNA EST 08-JUN-2000
DEFINITION hK49d05.y1 NCI_CGAP_Ov34 Homo sapiens cDNA clone IMAGE:3000009 5',
mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE

ORGANISM

human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE

1 (bases 1 to 50)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
cDNA Library Preparation: David B. Kitzman, Ph.D.
cDNA Library Arrayed by: I.M.A.G.E. Consortium, LLNL

JOURNAL

Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
cDNA Library Preparation: David B. Kitzman, Ph.D.
cDNA Library Arrayed by: I.M.A.G.E. Consortium, LLNL

DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL, send email to:
 info@image.lnl.gov

Seq primer: -40RP from Glbco.

FEATURES

source

Location/Qualifiers

1..50

/organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:3000009"
 /clone_lib="NCL_CGAP_OV34"
 /sex="female"
 /tissue_type="borderline ovarian carcinoma"
 /dev_stage="adult"
 /lab_host="DH10B"
 /note="Organ: ovary; Vector: pAMP1; mRNA made from
 borderline ovarian carcinoma, cDNA made by oligo-dT
 priming. Directionally cloned. Size-selected on agarose
 gel, average insert size 500 bp. Primary library,
 non-amplified."

BASE COUNT 30 a 3 c 11 g 6 t

ORIGIN

Query Match 4.3%; Score 14; DB 10; Length 50;
 Best Local Similarity 100.0%; Pred. No. 2.2e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 236 aattgtaaaaaa 249

Db 27 AATTGTAAAAAAA 40

RESULT 15

LOCUS

TA154F10Q 50 bp DNA GSS 13-DEC-2000

DEFINITION T. brucei sheared genomic DNA clone 154f10, reverse sequence,
 genomic survey sequence.

ACCESSION AL473287.1 GI:11838560

VERSION 1

KEYWORDS Trypanosoma brucei.

SOURCE Trypanosoma brucei

ORGANISM Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae;

Trypanosoma.

1 (bases 1 to 50)

AUTHORS

Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
 Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
 Melville, S.E., Rajadream, M.A. and Barrell, B.G.

Direct Submission

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
 project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
 Cambridgeshire CB10 1SA, E-mail: barrell@sanger.ac.uk and
 nh@sanger.ac.uk

COMMENT

Constructed at the Institute for Genomic Research (TIGR),
 Rockville, MD. Genomic DNA isolated from a cloned population of
 Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared
 to give a light size distribution (4 kb). The v + i method used for the library construction is
 described in detail in Smith, H. and Venter, J.C. (Making small
 insert libraries for whole genome shotgun sequencing projects. In
 Genome Sequencing: A Practical Approach, eds. M. Vaubin and B.
 Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org
 Details of T. brucei sequencing at the Sanger Centre are available
 at http://www.sanger.ac.uk/Projects/T_brucei/.

FEATURES

source

Location/Qualifiers

1..50

/organism="Trypanosoma brucei"
 /strain="TREU927"
 /db_xref="taxon:5691"
 /clone="154f10"

BASE COUNT

20 a 7 c 9 g 14 t

ORIGIN

Query Match 4.3%; Score 14; DB 13; Length 50;
 Best Local Similarity 100.0%; Pred. No. 2.2e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 268 atgaaaaaaatat 281

Db 10 ATGAAAAAATTTAT 23

RESULT 16

LOCUS

AZ331628 19 bp DNA GSS 29-SEP-2000

DEFINITION IM0059M12R Mouse 10kb plasmid UGCM1 library Mus musculus genomic
 clone UGCM1M0059M12 R, DNA sequence.

ACCESSION AZ331628

VERSION AZ331628.1 GI:10394503

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE

AUTHORS

1 (bases 1 to 19)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
 and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

Unpublished (2000)

JOURNAL

COMMENT

Contact: Robert B. Weiss
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177

Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0059 row: M column: 12
 Seq primer: CACACAGGAAACAGCTATGACC

Class: plasmid ends

High quality sequence stop: 19.

Location/Qualifiers

1..19

FEATURES

source

Location/Qualifiers

1..19

/organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UGCM1M0059M12"
 /clone_lib="Mouse 10kb plasmid UGCM1 library"
 /sex="male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (<http://www.jax.org/resources/documents/dnares/>). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 digested DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pMD42 (g114732114[gb]AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

```

BASE COUNT      8 a      1 c      0 g      10 t
ORIGIN
Query Match
Best Local Similarity 4.0%; Score 13; DB 13; Length 19;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 166 aaataaattag 178
Db 14 AAATAAATTAG 2

RESULT 17
AZ829725 25 bp DNA GSS 20-FEB-2001
LOCUS 2M0107124F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION clone UUGC2M0107124 F, DNA sequence.
ACCESSION AZ829725
VERSION AZ829725.1 GI:1299549
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 25)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0107 row: I column: 24
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 25.
Location/Qualifiers
1. 25
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0107124"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (9114732114|9b|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells

```

```

BASE COUNT      14 a      3 c      4 g      4 t
ORIGIN
Query Match
Best Local Similarity 4.0%; Score 13; DB 13; Length 25;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 301 ttatgtttcat 313
Db 24 TTTATGTTTCAT 12

RESULT 18
AZ309204 26 bp DNA GSS 29-SEP-2000
LOCUS 1M0013F07F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION clone UUGC1M0013F07 F, DNA sequence.
ACCESSION AZ309204
VERSION AZ309204.1 GI:10349955
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 26)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0013 row: F column: 07
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 26.
Location/Qualifiers
1. 26
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0013F07"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (9114732114|9b|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into

```

chemically-competent *E. coli* XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 4 a 4 c 4 g 14 t

ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttt 208
|||||

DB 7 TTATTTTATTTT 19

RESULT 19
A2866662/c

LOCUS 26 bp DNA GSS 21-FEB-2001
DEFINITION 2M0177A18F Mouse 10kb plasmid UGCG1M library Mus musculus genomic
clone UGCG2M0177A18 F, DNA sequence.

ACCESSION A2866662
VERSION A2866662.1 GI:13068193

KEYWORDS GSS.
SOURCE house mouse.

ORGANISM

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 26)

AUTHORS

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts
unpublished (2000)

JOURNAL

COMMENT Contact: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert length: 10000 Std Error: 0.00

Plate: 0177 row: A column: 18

Seq primer: CGTGTAAACGACGCCACGT

Class: plasmid ends

High quality sequence stop: 26.

FEATURES

SOURCE

1..26

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UGCG2M0177A18"

/clone_id="Mouse 10kb plasmid UGCG1M library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(<http://www.jax.org/resources/documents/dnares/>). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of PMD42 (g114732114[gb]AF129072.1), a copy-number

inducible derivative of plasmid RI. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adapted mouse DNA was annealed to

adapted vector DNA, and transformed into
chemically-competent *E. coli* XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 20 a 0 c 0 g 6 t

ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttt 208
|||||

DB 17 TTATTTTATTTT 5

RESULT 20
TA123B120/c

LOCUS 26 bp DNA GSS 13-DEC-2000
DEFINITION T. brucei sheared genomic DNA clone 123b12, reverse sequence,
genomic survey sequence.

ACCESSION AL463522
VERSION AL463522.1 GI:11834032

KEYWORDS GSS.

SOURCE Trypanosoma brucei.

ORGANISM Trypanosoma brucei.

Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;

Trypanosoma.

1 (bases 1 to 26)

Hall,N., Bowman,S., Leonard,N.J., Doggett,J., Atkin,R.,
Chillingworth,C., Ormond,D., Harris,B., El-Sayed,N., Hou,L.,
Melville,S.E., Rajandream,M.A. and Barrell,B.G.

Direct Submission

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing

project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,

Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and

nh@sanger.ac.uk

Constructed at the Institute for Genomic Research (TIGR),

Rockville, MD. Genomic DNA isolated from a cloned population of

Trypanosoma brucei (TREU927/4 G9at 10.1) was mechanically sheared

to give a tight size distribution (

4 kb). The v + i method used for the library construction is

described in detail in Smith, H. and Venter, J.C. (Making small

insert libraries for whole genome shotgun sequencing projects. In

Genome Sequencing: A Practical Approach, eds. M. Vaubin and B.

Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org

Details of T. brucei sequencing at the Sanger Centre are available

at http://www.sanger.ac.uk/Projects/T_brucei/.

FEATURES

SOURCE

1..26

/organism="Trypanosoma brucei"

/strain="TREU927"

/db_xref="taxon:5691"

/clone="123b12"

BASE COUNT 12 a 0 c 0 g 14 t

ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 271 aaaaaattattt 283
|||||

DB 20 AAAAAATTATTT 8

RESULT 21
A2784620

LOCUS 27 bp DNA GSS 16-FEB-2001
DEFINITION 2M0027P04R Mouse 10kb plasmid UGCG1M library Mus musculus genomic
clone UGCG2M0027P04 R, DNA sequence.

ACCESSION A2784620
 VERSION A2784620.1 GI:12920544
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 27)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamli,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0027 row: P column: 04
 Seq primer: CACACAGGAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 27.
 FEATURES
 source
 1..27
 Location/Qualifiers
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="U0GC2M0027P04"
 /clone_lib="Mouse 10kb plasmid U0GC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
 BASE COUNT
 5 a 5 c 5 g 12 t
 ORIGIN
 Query Match 4.0%; Score 13; DB 13; Length 27;
 Best Local Similarity 100.0%; Pred. No. 6.8e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 137 aaacttttttgt 149
 ||||||||||||
 Db 15 ACACTTTTGTGT 27
 RESULT 22
 A2452653 28 bp DNA GSS 04-OCT-2000
 LOCUS A2452653
 DEFINITION 1M0252E07R Mouse 10kb plasmid U0GC1M library Mus musculus genomic

clone U0GC1M0252E07 R, DNA sequence.
 ACCESSION A2452653
 VERSION A2452653.1 GI:10609676
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 28)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamli,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0252 row: E column: 07
 Seq primer: CACACAGGAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 28.
 FEATURES
 source
 1..28
 Location/Qualifiers
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="U0GC1M0252E07"
 /clone_lib="Mouse 10kb plasmid U0GC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
 BASE COUNT
 14 a 2 c 2 g 10 t
 ORIGIN
 Query Match 4.0%; Score 13; DB 13; Length 28;
 Best Local Similarity 100.0%; Pred. No. 6.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 271 aaaaaaatattt 283
 ||||||||||||
 Db 16 AAAAAATATTAT 28
 RESULT 23
 A2623794 30 bp DNA GSS 13-DEC-2000
 LOCUS A2623794/C

DEFINITION	1M0461C19R Mouse 10kb plasmid U08C1M library Mus musculus genomic clone U08C1M0461C19 R, DNA sequence.
ACCESSION	A2623794
VERSION	A2623794.1
KEYWORDS	GI:11745984
SOURCE	GSS.
ORGANISM	house mouse. Mus musculus.
REFERENCE	Eukaryotes. Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Scuriognathl; Muridae; Murinae; Mus. 1 (bases 1 to 30)
AUTHORS	Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A. and Wright, D., Weiss, R.
TITLE	Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL	Unpublished (2000)
COMMENT	Contact: Robert B. Weiss University of Utah Genome Center University of Utah Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA Tel: 801 585 5606 Fax: 801 585 7177 Email: addnegenetics.utah.edu Insert length: 10000 Std Error: 0.00 Plate: 0461 row: C column: 19 Seq primer: CACACAGGAACACGCTATGACC Class: plasmid ends High quality sequence stop: 30. location/Qualifiers 1..30 /organism="Mus musculus" /strain="C57BL/6J" /db_xref="taxon:10090" /clone="U08C1M0461C19" /clone_lib="Mouse 10kb plasmid U08C1M library" /sex="Male" /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-" /note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 (g1147311419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
BASE COUNT	14 a 4 c 3 g 9 t
ORIGIN	
Query Match	4.0%; Score 13; DB 13; Length 30;
Best Local Similarity	100.0%; Pred. No. 6.5e+05;
Matches	13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY	202 ttattttaattt 214
Db	14 TTATTTAAATTT 2

LOCUS	H40874	32 bp	mRNA		EST	31-JUL-1995
DEFINITION	y97ze10 s1Soares adult brain N2B5H855Y Homo sapiens cDNA clone IMAGE:1176394 3' similar to gb:X54150.ra1 IMMUNOGLOBULIN ALPHA FC RECEPTOR PRECURSOR (HUMAN); , mRNA sequence.					
ACCESSION	H40874					
VERSION	H40874.1	GI:916926				
KEYWORDS	EST.					
SOURCE	human.					
ORGANISM	Homo sapiens					
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.					
AUTHORS	1 (bases 1 to 32) Hallier,L., Clark,N., Dubouq,T., Elliston,K., Hawkin,M., Holman Ritkin,I., Rohlfing,T., Soares,M., Tan,F., Trevasakis,E., Waterston M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J. R., Williamson,A., Woldmann,P. and Wilson,R. The WashU-Merck EST Project Unpublished (1995)					
TITLE	Contact: Wilson RK					
JOURNAL	Washington University School of Medicine					
COMMENT	4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu Insert Size: 674 High quality sequence starts: 1 High quality sequence stops: 1 Source: IMAGE Consortium, LNL This clone is available royalty-free through LNL ; contact the IMAGE Consortium (infoimage.lnl.gov) for further information. Trace considered overall poor quality Insert Length: 674 Std Error: 0.00 Seq primer: Promega -21ml3 High quality sequence stop: 1. Location/Qualifiers 1..32 /organism="Homo sapiens" /db_xref="CDB:3838590" /db_xref="taxon:9606" /clone="IMAGE:1176394" /clone_1lb="Soares adult brain N2B5H855Y" /sex="Male" /dev_strage="55-year old" /lab_host="DH10B (ampicillin resistant)" /note="Organ: brain; Vector: pT73D (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5 TGTACCACCTGAAAGTGCGAGCGCCGGCTTTTATTTTTTTTTTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library went through one round of normalization to a Cot = 53. Library constructed by Bento Soares and M.Fatima Bonaldo. The adult brain RNA was provided by Dr. Donald H. Gilden. Tissue was acquired 17-18 hours after death which occurred in consequence of a ruptured aortic aneurysm. RNA was prepared from a pool of tissues representing the following areas of the brain: frontal, parietal, temporal and occipital cortex from the left and right hemispheres, subcortical white matter, basal ganglia, thalamus, cerebellum, midbrain, pons and medulla."					
BASE COUNT	6 a	0 c	0 g	26 t		
ORIGIN						
Query Match	4.0%; Score 13; DB 11; Length 32;					
Best Local Similarity	100.0%; Prid. No. 6.4e+05;					
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;						
196 ttattttatctt 208						
15 TTATTATTATTT 27						

```

RESULT 25
LOCUS      AZ458690
DEFINITION 1M0263E04F Mouse 10kb plasmid UGCC1M library Mus musculus genomic
ACCESSION  AZ458690
VERSION     AZ458690
KEYWORDS    GSS.
SOURCE      house mouse.
ORGANISM    Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE   1 (bases 1 to 32)
AUTHORS     Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
            Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
            ,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
            and Wright,D., Weiss,R.
TITLE       Mouse whole genome scaffolding with paired end reads from 10kb
            plasmid inserts
JOURNAL     Unpublished (2000)
COMMENT     Contact: Robert B. Weiss
            University of Utah Genome Center
            University of Utah
            Rm. 306, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
            84112, USA
            Tel: 801 585 5606
            Fax: 801 585 7177
            Email: ddunn@genetics.utah.edu
            Insert Length: 10000 Std Error: 0.00
            Plate: 0263 row: E column: 04
            Seq primer: CGTTGTAAACGACGCGCCAGT
            Class: plasmid ends
            High quality sequence stop: 32.
FEATURES
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        1..32
            Location/Qualifiers
                /organism="Mus musculus"
                /strain="C57BL/6J"
                /db_xref="taxon:10090"
                /clone="UUGC1M0263E04"
                /clone_1ib="Mouse 10kb plasmid UGCC1M library"
                /sex="Male"
                /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
                /note="Vector: PWD42nv; Purified genomic DNA from M.
                musculus C57BL/6J (male) was obtained from the Jackson
                Laboratory Mouse DNA Resource
                (http://www.jax.org/resources/documents/dnares/). The DNA
                was hydrodynamically sheared by repeated passage through a
                0.005 inch orifice at constant velocity. The sheared DNA
                was blunt end-repaired with T4 DNA polymerase and T4
                polynucleotide kinase. Adaptor oligonucleotides were
                ligated to the blunt ends in high molar excess. The
                adaptor DNA was purified and size-selected for a 9.5 to
                10.5 kb range using preparative agarose gel
                electrophoresis. Vector DNA was prepared from a derivative
                of PWD42 (q11473214|9b|AF129072.1), a copy-number
                inducible derivative of plasmid RL. The vector was ligated
                with adaptors complementary to the insert adaptors and
                purified. The sheared, adaptor mouse DNA was annealed to
                adaptor vector DNA, and transformed into
                chemically-competent E. coli XL10-Gold (Stratagene) cells
                and selected for ampicillin resistance."
BASE COUNT      4 a
ORIGIN           5 c
Query Match      4.0%; Score 13; DB 13; Length 32;
Best Local Similarity 100.0%; Pred. No. 6.4e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 196 ttaatttttttt 208
|||||

```

```

Db 12 TTATTTTATTTT 24

RESULT 26
LOCUS      AA906810/C
DEFINITION ok73c02.s1 NCI_CGAP_GC4 Homo sapiens cDNA clone IMAGE:1519586
ACCESSION  AA906810
VERSION     AA906810.1 GI:3042054
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE   1 (bases 1 to 34)
AUTHORS     NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE       National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL     Unpublished (1997)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
            Emmert-Buck, M.D., Ph.D.
            CDNA Library Preparation by: Bento Soares, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/bbrp/image/image.html
            Trace considered overall poor quality
            Seq primer: -40m13 fwd. ET from Amerisham
            High quality sequence stop: 1.
FEATURES
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        1..34
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                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="IMAGE:1519586"
                /clone_1ib="NCI_CGAP_GC4"
                /tissue_type="pooled germ cell tumors"
                /lab_host="DH10B"
                /note="Vector: pTZ19-Pac (Pharmacia) with a modified
                polylinker: 1st strand cDNA was prepared from 3 pooled
                germ cell tumors, and was then primed with a Not I -
                oligo(dT) primer. Double-stranded cDNA was ligated to Eco
                RI adaptors (Pharmacia), digested with Not I and cloned
                into the Not I and Eco RI sites of the modified pTZ19
                vector. Library is normalized. Library was constructed by
                Bento Soares and M. Fatima Bonaldo."
BASE COUNT      8 a
ORIGIN           4 c
Query Match      4.0%; Score 13; DB 10; Length 34;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 51 ataataaagaat 63
|||||
Db 21 ATATATATAAAGAT 9

RESULT 27
LOCUS      AZ586746
DEFINITION 1M0392K22R Mouse 10kb plasmid UGCC1M library Mus musculus genomic
ACCESSION  AZ586746
VERSION     AZ586746.1 GI:11708936
KEYWORDS    GSS.
SOURCE      house mouse.

```


ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 34)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0392 row: K column: 22
Seq primer: CACACAGCAACACGTATGACC
Class: plasmid ends
High quality sequence stop: 34.
Location/Qualifiers
1. 34
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C1M0392K22"
/clone_lib="Mouse 10kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gblAF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 2 a 0 c 4 g 28 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 34;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 196 ttattttatttt 208
|||||
Db 6 TTATTTTATTTT 18

RESULT 28
A2781725 34 bp DNA GSS 16-FEB-2001
LOCUS
DEFINITION 2M002111AF Mouse 10kb plasmid U06C1M library Mus musculus genomic
ACCESSION A2781725
VERSION A2781725.1 GI:12914706
KEYWORDS GSS.

SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 34)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0021 row: I column: 14
Seq primer: CGTGTAAACGACGCGCAGT
Class: plasmid ends
High quality sequence stop: 34.
Location/Qualifiers
1. 34
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C2M0021I14"
/clone_lib="Mouse 10kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gblAF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 13 a 4 c 4 g 13 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 34;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 209 aaattgttaaaa 221
|||||
Db 10 AAATTGTTAAAA 22

RESULT 29
AM246486 35 bp mRNA EST 07-JAN-2000
LOCUS
DEFINITION 2821545.3prime NIH_MGC_7 Homo sapiens cDNA clone IMAGE:2821545 3',
ACCESSION AM246486
VERSION AM246486.1 GI:6589479

KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
TITLE 1 (bases 1 to 35)
JOURNAL NIH-MGC http://mgi.nci.nih.gov/.
COMMENT National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Other_ESTS: 2821545.5prime
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: DCTD/DPB CDNA Library Preparation: Lung
Hong/Rubin Laboratory CDNA Library Arrayed by: The I.M.A.G.E.
Consortium (LNL) DNA Sequencing by: Berkeley MGC sequencing
project Clone distribution: MGC clone distribution information can
be found through the I.M.A.G.E. Consortium/LNL at:
www.bio.lnl.gov/dbp/image/image.html Base Calling / Quality
Scores: PHRED from University of Washington Genome Center
Trimming: crossmatch from University of Washington Genome Center
PHRAP suite. Poly-T identification: patmatch.pl from Berkeley
Drosophila Genome Project. University of Washington Genome Center:
http://www.genome.washington.edu Low Quality Sequence: 35
contiguous PHRED high quality bases following vector sequence. Very
Low Quality Sequence: Trace file contained 35 contiguous distinct
peaks following vector sequence. Polyadenylation: Based upon the
presence of a XhoI site followed by a run of 14 or more T residues
at the beginning of the sequence, this CDNA insert was
polyadenylated.
Plate: LCM7 row: B column: 10
High quality sequence stop: 35.
Location/Qualifiers
1..35
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="NIH-MGC-7"
/tissue_type="small cell carcinoma"
/cell_line="MGC3"
/lab_host="DH10B (phage-resistant)"
/note="Organ: Lung; Vector: pOT7; Site_1: XhoI; Site_2:
EcoRI; CDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT 11 a 4 c 1 g 19 t
ORIGIN
Query Match 4.0%; Score 13; DB 10; Length 35;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 35 tttaaatgaata 47
|||||
Db 8 TTTAAATGAATA 20

RESULT 30
LOCUS D45807 35 bp mRNA EST 20-FEB-1995
DEFINITION HKMS03025 Human adult lung 3' directed MboI CDNA Homo sapiens CDNA
3', mRNA sequence.
ACCESSION D45807
VERSION D45807.1 GI:662761
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 35)
AUTHORS Itoh, K., Okubo, K., Yosi, J., Yokouchi, H. and Matsubara, K.
TITLE An expression profile of active genes in human lung
JOURNAL DNA Research 1, 279-287 (1994)
MEDLINE 95236275
COMMENT Contact: Kohichi Itoh
Institute for Molecular and Cellular Biology
Osaka University
3-1, Yamadaoka, Suita, Osaka, 565, Japan
Tel: 06-877-5111 x3910
Fax: 06-877-1922.
FEATURES
source
1..35
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Human adult lung 3' directed MboI CDNA"
/note="Adult human lung, 3' directed MboI"
BASE COUNT 14 a 3 c 5 g 13 t
ORIGIN
Query Match 4.0%; Score 13; DB 11; Length 35;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 106 atagtttaaat 118
|||||
Db 11 ATAGTTTAATAAT 23

RESULT 31
LOCUS AZ314238 36 bp DNA GSS 29-SEP-2000
DEFINITION 1M0030N24R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0030N24 R, DNA sequence.
ACCESSION AZ314238
VERSION AZ314238.1 GI:10359929
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 36)
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacons, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Rilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A.
and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0030 row: N column: 24
Seq primer: CACACGAGAACGCTATGACC
Class: plasmid ends
High quality sequence stop: 36.
Location/Qualifiers
1..36
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone_lib="UUGC1M0030N24"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g14732114[gb]/AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent *E. coli* XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 8 a 3 c 5 g 20 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 36;
Best Local Similarity 100.0%; Pred. No. 6;le+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 312 atgtttcttatt 324
|||||
Db 2 ATGTTTCTATT 14

RESULT 32
LOCUS AA913140 37 bp mRNA 14-APR-1998
DEFINITION o118h06.s1 NCI-CGAP_HNI Homo sapiens cDNA clone IMAGE:1483067 3'
similar to SW:YK13_YEAST P36079 HYPOTHETICAL 23.7 KD PROTEIN IN
MDM1-VMA5 INTERGENIC REGION. ;, mRNA sequence.
ACCESSION AA913140
VERSION AA913140.1 GI:3052532
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 37)
NCI/NIH-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute / National Institute of Dental Research,
Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
Unpublished (1997)
CONTACT: Robert Strausberg, Ph.D.
Email: cgapbs-rt@mail.nih.gov
Tissue Procurement: John Ensley, M.D., Mary May, J. Silvio Gutkind,
Ph.D.

CDNA Library Preparation: Stratagene, Inc.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
www-bio.llnl.gov/bdrip/image/image.html

Trace considered overall poor quality
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 1.

FEATURES

Source
1. .37
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1483067"
/clone_lib="NCI-CGAP_HNI"
/issue_type="squamous cell carcinoma"
/lab_host="SOLR (kanamycin resistant)"
/note="Organ: lymph node; Vector: Bluescript SK-; Site_1:
EcoRI; Site_2: XhoI; Cloned unidirectionally. Primer:

OLigo dT. Average insert size 1.3 kb. 5' adaptor sequence:
5' GAATTCGGCAGCAG 3' 3' adaptor sequence: 5' (GA
1)GACATGCTCGAGCTTTTCTTTTCTTTT 3' "

BASE COUNT 2 a 9 c 2 g 24 t
ORIGIN

Query Match 4.0%; Score 13; DB 10; Length 37;
Best Local Similarity 100.0%; Pred. No. 6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 196 ttattttatttt 208
|||||
Db 3 TTATTTTATTATT 15

RESULT 33
LOCUS AU009123 37 bp mRNA 31-JUL-1998
DEFINITION AU009123 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA clone spc04510, mRNA sequence.
ACCESSION AU009123
VERSION AU009123.1 GI:3345803
KEYWORDS EST.
SOURCE fission yeast.
ORGANISM Schizosaccharomyces pombe

Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetaceae; Schizosaccharomycetaceae;
Schizosaccharomycetes.
1 (bases 1 to 37)
Moriyomo,M. and Mita,K.
Identification of expressed sequence tags of Schizosaccharomyces
pombe

Unpublished (1998)
CONTACT: MitsuoKI Moriyomo
Genome Research Group
National Institute of Radiological Sciences
9-1, Anagawa 4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
Email: moriyomo@nirs.go.jp.
Location/Qualifiers

FEATURES

Source
1. .37
/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc04510"
/clone_lib="Schizosaccharomyces pombe late log phase cDNA"
/sex="h minus"
/note="Vector: M13mp19; The cDNA library of
Schizosaccharomyces pombe was prepared by cloning cDNA
into the SmaI site of M13mp19 DNA and the direction of DNA
sequences was not always from 5' to 3'. The cDNA data of
Schizosaccharomyces pombe are available for searching on
the World Wide Web. (URL, <http://www.nirs.go.jp>)"

BASE COUNT 13 a 6 c 3 g 15 t
ORIGIN

Query Match 4.0%; Score 13; DB 10; Length 37;
Best Local Similarity 100.0%; Pred. No. 6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 110 ttataataaaaa 122
|||||
Db 13 TTAATTAATAAAA 25

RESULT 34
LOCUS AA916625 40 bp mRNA 10-JUN-1998
DEFINITION om05g12.s1 Soares_NFL_T_GRC_S1 Homo sapiens cDNA clone
IMAGE:1540198 3 similar to SW:NO5M.TRYBB P04540 NADH-UBIQUINONE
OXIDOREDUCTASE CHAIN 5 ;, mRNA sequence.
ACCESSION AA916625

```

VERSION AA916625.1 GI:3056017
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Euthera; Primates; Catarrhini; Homidae; Homo.
TITLE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
JOURNAL National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
COMMENT Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-rt@mail.nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 1026 Std Error: 0.00
Seq primer: -40m3 fwd. ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
1. 40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_image="1540198"
/clone_lib="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site: 1: Not I; Site: 2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NBH13W, testis NHT, and B-cell
NCI-CGAP-CGI) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 662632-667239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo. "
BASE COUNT 10 a 1 c 2 g 27 t
ORIGIN
Query Match 4.0%; Score 13; DB 10; Length 40;
Best Local Similarity 100.0%; Pred. No. 5.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 106 atagtaataaat 118
|||||
Db 21 ATAGTAATAAAT 9

RESULT 35
AA922076/c 40 bp mRNA EST 21-APR-1998
LOCUS oh08907.s1 NCI-CGAP Co8 Homo sapiens cDNA clone IMAGE:1457244 3'
DEFINITION similar to TR:Q34096 Q34096 MURF2 PROTEIN. ; mRNA sequence.
ACCESSION AA922076
VERSION AA922076.1 GI:3069385
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Euthera; Primates; Catarrhini; Homidae; Homo.
TITLE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
JOURNAL National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
COMMENT Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-rt@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.

```

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cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/btrp/image/image.html
Trace considered overall poor quality
Seq primer: -40m3 fwd. ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
1. 40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_image="1457244"
/clone_lib="NCI CGAP Co8"
/tissue_type="adenocarcinoma"
/lab_host="DH10B"
/note="Organ: colon; Vector: pT7T3D-Pac (Pharmacia) with a
modified polylinker; 1st strand cDNA was prepared from
colon adenocarcinoma, and was then primed with a Not I -
oligo(drf) primer. Double-stranded cDNA was ligated to Eco
RI adaptors (Pharmacia), digested with Not I and cloned
into the Not I and Eco RI sites of the modified pT7T3
vector. Library is normalized. Library was constructed by
Bento Soares and M. Fatima Bonaldo. "
BASE COUNT 5 a 4 c 3 g 28 t
ORIGIN
Query Match 4.0%; Score 13; DB 10; Length 40;
Best Local Similarity 100.0%; Pred. No. 5.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 112 aatataataaag 124
|||||
Db 38 AATATAATAAAG 26

RESULT 36
C00434 41 bp mRNA EST 23-JUL-1996
LOCUS HUMGS0006099 Human adult (K.Okubo) Homo sapiens cDNA, mRNA
DEFINITION sequence.
ACCESSION C00434
VERSION C00434.1 GI:1432664
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Euthera; Primates; Catarrhini; Homidae; Homo.
TITLE Okubo, K.
JOURNAL BodyMap: human gene expression database
COMMENT Unpublished (1995)
Contact: Okubo, K.
Institute for Molecular and Cellular Biol
Osaka University
1-3 Yamada-oka, Suita, Osaka Pref. 565, Japan
Tel: 06-877-5111(ex.3315)
Email: kousaku@imcb.osaka-u.ac.jp
Human Gene Signature, 3'-directed cDNA sequence. We are not
submitting the same cDNA sequence redundantly to DBJ since 1993.
For the abundance information of clones with this sequence in this
library and as well as in other 3'-directed libraries, see
http://www.imcb.osaka-u.ac.jp/bodymap/. The sequences of the clones
represented by this GS sequences is also found there.
Location/Qualifiers
1. 41
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Human adult (K.Okubo)"
/dev_stage="adult"
BASE COUNT 12 a 5 c 7 g 17 t

```

ORIGIN

Query Match 4.0%; Score 13; DB 11; Length 41;
 Best Local Similarity 100.0%; Pred. No. 5.8e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 109 gftataataaa 121
 |||||||||||||
 Db 14 GTTAATAATAA 26

RESULT 37

A2777050/c

LOCUS A2777050 41 bp DNA GSS 16-FEB-2001
 DEFINITION 2M001M12F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC2M001M12 F, DNA sequence.

ACCESSION

A2777050

A2777050.1 GI:12905260

VERSION

KEYWORDS

SOURCE

ORGANISM

house mouse.

Mus musculus

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Contact: Robert B. Weiss

University of Utah

Rm. 308, Biomedical

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000

Std Error: 0.00

Plate: 0011 row: M

Column: 12

Seq primer: CGTGTAAACGACGCCAGT

Class: plasmid ends

High quality sequence stop: 41.

Location/Qualifiers

1. 41

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC2M001M12"

/clone_lib="Mouse 10kb plasmid UUGC1M library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of PMD42 (g14732114[gblAF129072.1]), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adapted mouse DNA was annealed to

chemically-competent E. coli XL10-Gold (Stratagene) cells

and selected for ampicillin resistance."

BASE COUNT

9 a 3 c 12 g 17 t

ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 41;
 Best Local Similarity 100.0%; Pred. No. 5.8e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 57 aaaagatatata 69
 |||||||||||||
 Db 16 AAAAGATATATTA 4

RESULT 38

A2371136

LOCUS A2371136 43 bp DNA GSS 02-OCT-2000
 DEFINITION 1M0122N05F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0122N05 F, DNA sequence.

ACCESSION

A2371136

A2371136.1 GI:10484836

VERSION

KEYWORDS

SOURCE

ORGANISM

house mouse.

Mus musculus

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Contact: Robert B. Weiss

University of Utah

Rm. 308, Biomedical

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000

Std Error: 0.00

Plate: 0122 row: N

Column: 05

Seq primer: CGTGTAAACGACGCCAGT

Class: plasmid ends

High quality sequence stop: 43.

Location/Qualifiers

1. 43

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC1M0122N05"

/clone_lib="Mouse 10kb plasmid UUGC1M library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of PMD42 (g14732114[gblAF129072.1]), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adapted mouse DNA was annealed to

chemically-competent E. coli XL10-Gold (Stratagene) cells

and selected for ampicillin resistance."

BASE COUNT 28 a 2 c 2 g 11 t
 and selected for ampicillin resistance."

Query Match 4.0%; Score 13; DB 13; Length 43;
 Best Local Similarity 100.0%; Pred. No. 5.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 111 taataataaaaa 123
 |||
 Db 18 TAATAATAAAAA 30

RESULT 39

AZ464392 43 bp DNA GSS 04-OCT-2000
 LOCUS 1M0273D17R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 DEFINITION clone UUGC1M0273D17 R, DNA sequence.

ACCESSION AZ464392
 VERSION AZ464392.1 GI:10622517
 KEYWORDS GSS.

SOURCE house mouse.
 ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 43)

AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Haml, C.,
 Islam, H., Longacre, S., Mahmood, M., Meenen, E., Pedersen, T., Reilly,
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A.
 and Wright, D., Weiss, R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA

Tel: 801 585 5606
 Fax: 801 585 7177

Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0273 row: D column: 17

Seq primer: CACACAGAAACAGCATGACC
 Class: plasmid ends

High quality sequence stop: 43.
 Location/Qualifiers

1..43
 /organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC1M0273D17"

/clone_lib="Mouse 10kb plasmid UUGC1M library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PMD42ny: Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 ligated DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of PMD42 (g114732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptor complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into

BASE COUNT 6 a 7 c 8 g 22 t
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

Query Match 4.0%; Score 13; DB 13; Length 43;
 Best Local Similarity 100.0%; Pred. No. 5.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 139 accttttgcctt 151
 |||
 Db 30 ACTTTTGTGTTT 42

RESULT 40

AZ575514 43 bp DNA GSS 06-DEC-2000
 LOCUS AZ575514
 DEFINITION AST-T21F0045 Genetrap T47D Human Breast Carcinoma Library Homo
 sapiens genomic 5', DNA sequence.

ACCESSION AZ575514
 VERSION AZ575514.1 GI:11561825
 KEYWORDS GSS.

SOURCE human.
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 1 (bases 1 to 43)

AUTHORS Henkel, G., Liyanage, M., Pratt, E., Huang, D., Riley, M., Bernardino, A.,
 Durick, K., and Pollok, B.

TITLE Exon-trap tags from a T47D Genomescreen(TM) Library
 JOURNAL Unpublished (2000)

COMMENT Contact: Greg Henkel
 Gene Expression
 Aurora Biosciences Corp.
 11010 Torreyana Road, San Diego, CA 92121, USA
 Tel: 8584048436
 Fax: 8584046719

Email: henkelg@aurora.bio.com

Pools of cells were isolated from a Genomescreen(TM) library. The
 library of cells was generated by retroviral integration of a gene
 tagging element consisting of: 1) A promoterless beta-lactamase
 proceeded by a splice acceptor as a reporter for gene expression;
 2) A promoter driving neomycin resistance followed by a splice
 donor to trap downstream exons. 3' RACE from neomycin gene was
 performed using total RNA from isolated pools. Output was shotgun
 cloned in pAMP-1 and used to transform DH5-alpha competent
 bacteria. 5' ends of reported sequences were immediately preceded
 by splice donor from the trapping construct.
 Class: exon-trapped.

Location/Qualifiers

1..43
 /organism="Homo sapiens"

/db_xref="taxon:9606"

/clone_lib="Genetrap T47D Human Breast Carcinoma Library"

/tissue_type="Carcinoma"

/cell_type="Epithelial"

/cell_line="T47D"

/note="Organ: Breast; Vector: pAMP-1; 3' RACE of total RNA
 from genetrap pools; shotgun clone in pAMP-1 and used to
 transform DH5-alpha competent bacteria."

BASE COUNT 18 a 9 c 5 g 11 t

Query Match 4.0%; Score 13; DB 13; Length 43;
 Best Local Similarity 100.0%; Pred. No. 5.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 aatgaataaaga 51
 |||

Db 1 AATGAATAAAGA 13

RESULT 41
D20668/c 45 bp mRNA EST 30-JUL-1996
LOCUS D20668 Human promyelocyte Homo sapiens cDNA clone pm2268 3',
DEFINITION HMG501644 Human promyelocyte Homo sapiens cDNA clone pm2268 3',
RNA sequence.
ACCESSION D20668
VERSION D20668.1 GI:501764
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
AUTHORS Okubo, K., Fukushima, A., Yoshi, J., Niyama, T., Kojima, Y., Yoshinari,
H., Arimoto, J., and Matsubara, K.
TITLE Gene expression of human promyelocytic cell line HL60 before and
after induction of differentiation. A new application of 3'directed
cDNA sequencing
Unpublished (1993)
JOURNAL Contact: Okubo, K., Fukushima, A., Yoshi, J., Niyama, T., Kojima, Y.,
COMMENT Yoshihara, H., Arimoto, J., and Matsubara, K.
Institute for Molecular and Cellular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.
FEATURES
SOURCE Location/Qualifiers
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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="pm2268"
/clone_lib="Human promyelocyte"
/note="Female, adult, cell_line = HL60, cell_type =
promyelocyte."
BASE COUNT 17 a 6 c 6 g 14 t 2 others
ORIGIN

Query Match 4.0%; Score 13; DB 11; Length 45;
Best Local Similarity 100.0%; Pred. No. 5.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 196 ttattttatttt 208
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Db 34 TTTATTTTATTTT 22

RESULT 42
A2459612 46 bp DNA GSS 04-OCT-2000
LOCUS A2459612 Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION clone UUGC1M0264002 R, DNA sequence.
ACCESSION A2459612
VERSION A2459612.1 GI:10617737
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
JOURNAL Contact: Robert B. Weiss
COMMENT University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177

Email: ddunne@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0264 row: 0 column: 02
Seq primer: CACACAGCAACACGCTATGACC
Class: plasmid ends
High quality sequence stop: 46.
Location/Qualifiers
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/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0264002"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g11473211419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance.
BASE COUNT 21 a 5 c 4 g 16 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 46;
Best Local Similarity 100.0%; Pred. No. 5.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 190 aaataattatttt 202
|||||
Db 32 AAATAATTATTTT 44

RESULT 43
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LOCUS A2834972/c
DEFINITION 2M017N19R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M017N19 R, DNA sequence.
ACCESSION A2834972
VERSION A2834972.1 GI:13004880
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
JOURNAL Contact: Robert B. Weiss
COMMENT University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606

Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0117 row: N column: 19
 Seq primer: CACACAGCAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 46.

FEATURES

SOURCE

Location/Qualifiers

1. 46
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUCG2M0117N19"
 /clone_lib="Mouse 10kb plasmid UUCG1M library"
 /sex="Male"
 /lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT

15 a 7 c 9 g 15 t

BASE COUNT

23 a 10 c 5 g 8 t

Query Match 4.0%; Score 13; DB 13; Length 46;
 Best Local Similarity 100.0%; Pred. No. 5.6e+05;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 30 gaatgttaaatc 42
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 Db 20 GAATGTTAAATG 8

RESULT 44

AZ991460

LOCUS

DEFINITION 2M0275C07R Mouse 10kb plasmid UUCG2M library Mus musculus genomic
 clone UUCG2M0275K07 R, DNA sequence.

ACCESSION

AZ991460

VERSION

AZ991460.1

KEYWORDS

GSS.

SOURCE

house mouse.

ORGANISM

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 46)

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,

Islam,H., Longacre,S., Mahmood,M., Meenen,E., Pedersen,T., Reilly,

M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.

and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

JOURNAL

COMMENT

Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0275 row: K column: 07
 Seq primer: CACACAGCAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 46.

FEATURES

SOURCE

Location/Qualifiers

1. 46
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUCG2M0275K07"
 /clone_lib="Mouse 10kb plasmid UUCG2M library"
 /sex="Female"
 /lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (female) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

Query Match 4.0%; Score 13; DB 13; Length 46;
 Best Local Similarity 100.0%; Pred. No. 5.6e+05;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 taatgaataaa 49
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 Db 32 TAAATGAATAAA 44

RESULT 45

A1630064

LOCUS

DEFINITION A1630064 49 bp mRNA EST 08-MAR-2000
 cd00157 Proliferating Erythroid Cells (LCB-ad library) Homo sapiens
 cDNA clone ad00157 random, mRNA sequence.

ACCESSION

A1630064

VERSION

A1630064.1

KEYWORDS

EST.

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

1 (bases 1 to 49)

Gubin,A.N., Njoroge,J.M., Bouffard,G.G. and Miller,J.L.

Gene expression in proliferating human erythroid cells

Genomics 59 (2), 168-177 (1999)

9939981

Contact: Jeffery L. Miller

Laboratory of Chemical Biology

National Institute of Diabetes and Digestive and Kidney Diseases

Building 10, Room 9B17, National Institutes of Health, Bethesda, MD

20892, USA

Tel: 301 402 2373

Fax: 301 435 5148

JOURNAL

COMMENT

Email: jmf@nih.gov
 The 'ad' library was constructed by Alexander Gubin, Ph.D. in the
 Laboratory of Chemical Biology, NIDDK, NIH. DNA Sequencing and/or
 analyses by National Institutes of Health Intramural Sequencing
 Center (NISC). More information available at:
<http://hembase.nidck.nih.gov>.

FEATURES

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 /db_xref="taxon:9606"
 /clone="ad00157"
 /clone_lib="Proliferating Erythroid Cells (LCB:ad library)"
 /sex="unknown"
 /tissue_type="blood"
 /cell_type="Erythroid Cells"
 /cell_line="Primary Culture of Peripheral Blood Mononuclear Cells"
 /dev_stage="Progenitor; EPO responsive CD71+++"
 /lab_host="DH5alpha"
 /note="Organ: blood; Vector: pCRIT; Site_1: EcoRI; Site_2: EcoRI; Human peripheral blood mononuclear cells were cultured in the presence or absence of erythropoietin. CD71+++ cells arising only in erythropoietin-supplemented medium were isolated by fluorescence activated cell sorting. Those cells demonstrated an average of six additional doublings in suspension culture and erythroid colony formation in methylcellulose. Suppression subtractive hybridization was used to construct the ad library (tester-sorted CD71+++ cells, driver=unsorted cells cultured without erythropoietin)."
 BASE COUNT 16 a 7 c 9 g 13 t 4 others
 ORIGIN

Query Match 4.0%; Score 13; DB 10; Length 49;
 Best Local Similarity 100.0%; Pred. No. 5.4e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 172 aattagataaaa 184
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 Db 24 AATTGATGATAAAA 36

Search completed: January 24, 2002, 02:57:29
 Job time: 3951 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:13:27 ; Search time 90.4 Seconds

(without alignments)
819.229 Million cell updates/sec

Title: US-09-531-438-3

Perfect score: 327
Sequence: 1 atttgagatacttaattt.....tttcattgtttctattgtt 327

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 351203 seqs, 113238999 residues

Total number of hits satisfying chosen parameters: 702406

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

Issued Patents_NA: *
1: /cgn2.6/ptodata/2/ina/5A.COMB.seq: *
2: /cgn2.6/ptodata/2/ina/5B.COMB.seq: *
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4: /cgn2.6/ptodata/2/ina/6B.COMB.seq: *
5: /cgn2.6/ptodata/2/ina/PC105.COMB.seq: *
6: /cgn2.6/ptodata/2/ina/Backfile1.seq: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	63.8	19.5	2435	US-09-306-593-1	Sequence 1, Appli
2	62.2	19.0	6243	US-09-056-075-1	Sequence 1, Appli
3	61.8	18.9	615	US-08-998-416-186	Sequence 186, App
4	61.6	18.8	19124	US-08-487-826B-13	Sequence 13, Appli
5	59.2	18.1	5852	US-07-867-106-2	Sequence 2, Appli
6	58.2	17.8	9048	US-08-973-273-4	Sequence 4, Appli
7	57.8	17.7	636	US-08-998-416-1137	Sequence 1137, Ap
8	56.8	17.4	665	US-08-883-795A-36	Sequence 36, Appli
9	56.4	17.2	837	US-08-998-416-288	Sequence 288, App
10	55.8	17.1	2750	US-08-617-860B-33	Sequence 33, Appli
11	55.6	17.0	665	US-08-883-795A-36	Sequence 36, Appli
12	54.8	16.8	8920	US-08-446-855A-1	Sequence 1, Appli
13	54.8	16.8	8920	US-09-150-741-1	Sequence 1, Appli
14	54.6	16.7	2251	US-08-991-677-11	Sequence 11, Appli
15	54.4	16.6	5852	US-07-867-106-2	Sequence 2, Appli
16	53.4	16.3	658	US-08-998-416-595	Sequence 595, App
17	53.2	16.3	3095	US-08-1168-1	Sequence 1, Appli
18	52.8	16.1	2110	US-09-419-459-1	Sequence 1, Appli
19	52.8	16.1	3095	US-09-1168-1	Sequence 1, Appli
20	52.6	16.1	1826	US-09-286-691-11	Sequence 11, Appli
21	52.6	16.1	1826	US-09-687-147-11	Sequence 11, Appli
22	52.6	16.1	3305	US-09-068-043-1	Sequence 1, Appli
23	52.6	16.1	51952	US-08-947-823-1	Sequence 1, Appli
24	52.2	16.0	8920	US-08-446-855A-1	Sequence 1, Appli
25	52.2	16.0	8920	US-09-150-741-1	Sequence 1, Appli
26	52	15.9	19124	US-08-487-826B-13	Sequence 13, Appli
27	51.2	15.7	1186	US-08-731-722-5	Sequence 5, Appli

28	51.2	15.7	2230	US-08-378-313-24	Sequence 24, Appli
29	50.8	15.5	1477	US-08-096-181A-7	Sequence 7, Appli
30	50.8	15.5	1477	PCT-US94-08326-7	Sequence 7, Appli
31	50.6	15.5	860	US-08-998-416-287	Sequence 287, App
32	50.6	15.5	2110	US-09-419-459-1	Sequence 1, Appli
33	50.4	15.4	1431	US-09-316-083-2	Sequence 2, Appli
34	50.2	15.4	1850	US-08-617-860B-32	Sequence 32, Appli
35	50.2	15.4	4098	US-08-605-106-4	Sequence 4, Appli
36	50.2	15.3	660	US-07-991-867B-32	Sequence 32, Appli
37	50	15.3	660	US-08-107-755A-32	Sequence 32, Appli
38	50	15.3	660	US-08-544-332-32	Sequence 32, Appli
39	50	15.3	734	US-09-014-583-1	Sequence 1, Appli
40	50	15.3	1511	US-07-991-867B-8	Sequence 8, Appli
41	50	15.3	1511	US-08-107-755A-8	Sequence 8, Appli
42	50	15.3	1511	US-08-544-332-8	Sequence 8, Appli
43	50	15.3	1667	US-08-485-284A-1	Sequence 1, Appli
44	50	15.3	4810	US-08-852-629-11	Sequence 11, Appli
45	50	15.3	4838	US-08-852-629-15	Sequence 15, Appli

ALIGNMENTS

```
RESULT 1
US-09-306-593-1
; Sequence 1, Application US/09306593
; Patent No. 6184018
;
GENERAL INFORMATION:
; APPLICANT: Li, Xin-Liang
; APPLICANT: Ljungdahl, Lars G.
; APPLICANT: Chen, Huizhong
; APPLICANT: Ximenes, Eduardo A.
;
TITLE OF INVENTION: Beta-glucosidase Coding Sequences and Protein from
; FILE REFERENCE: 31-98us
;
CURRENT APPLICATION NUMBER: US/09/306,593
; CURRENT FILING DATE: 1999-05-06
; EARLIER APPLICATION NUMBER: US 60/084,494
; EARLIER FILING DATE: 1998-05-06
;
NUMBER OF SEQ ID NOS: 13
;
SOFTWARE: PatentIn Ver. 2.0
;
SEQ ID NO 1
; LENGTH: 2435
; TYPE: DNA
; ORGANISM: Orpinomyces sp. PC-2
;
FEATURE:
; NAME/KEY: CDS
; LOCATION: (39)..(2009)
; FEATURE:
; NAME/KEY: mat_peptide
; LOCATION: (87)..(2009)
;
US-09-306-593-1
;
Query Match
Best Local Similarity 56.9%; Score 63.8; DB 4; Length 2435;
Matches 136; Conservative 0; Mismatches 102; Indels 1; Gaps 1;
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OY 43 aaataaagataaataaagataataataatgctgaataatataaata-taagat 101
DB 2052 aaataaataataataaataatgataatatttatttactcttttgctaaagtaagaat 2111
OY 102 aagatagtaataataaataaagttctcgagggaactttttgttttaaaaggaa 161
DB 2112 aaataaatttataataataataatgataataatcttttgaatcattaaata 2171
OY 162 atataataaatttgaataaagttgaataatatttttaatttgaatttgaataa 221
DB 2172 aaataaataataaataaatttgaataataatgataatatttataatatttgaaga 2231
OY 222 attgataataatgataatgtaaaaaatttcoagggggaatataatgtaaaaaata 280
DB 2232 gattataatttataaataataataaagaagaacaaataataataataaaaaata 2290
```

RESULT 2

US-09-056-075-1

Sequence 1 Application US/09056075
Patent No. 5955368

GENERAL INFORMATION:

APPLICANT: Johnson, Eric A.
APPLICANT: Bradshaw, Marie
APPLICANT: Rood, Julian
TITLE OF INVENTION: Expression System for Clostridium
TITLE OF INVENTION: Species
NUMBER OF SEQUENCES: 2
CORRESPONDENCE ADDRESS:
ADDRESSEE: Quarles & Brady
STREET: 1 South Pluckney Street
CITY: Madison
STATE: WI
COUNTRY: US
ZIP: 53701-2113

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/056,075

FILING DATE:

CLASSIFICATION:

ATTORNEY/AGENT INFORMATION:
NAME: Seay, Nicholas J.
REGISTRATION NUMBER: 27386
REFERENCE/DOCKET NUMBER: 960296.95238
TELECOMMUNICATION INFORMATION:
TELEPHONE: 608-251-5000
TELEFAX: 608-251-9166

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:
LENGTH: 6243 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: misc.feature
LOCATION: 3770..4013
OTHER INFORMATION: /note="RP4 origin of DNA transfer (oriT) from
US-09-056-075-1
OTHER INFORMATION: plasmid RP4"

Query Match 19.0%; Score 62.2; DB 2; Length 6243;

Best Local Similarity 49.5%; Pred. No. 0.0071;
Matches 160; Conservative 0; Mismatches 163; Indels 0; Gaps 0;

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DB 1183 ATGACGGCTAAATTAATAATAATAATAATAATAATAATAATA 1242
QY 61 gatataatattatagctgaataattatataatgataagtaataa 120
DB 1243 AATTAATAATAATAATAATAATAATAATAATAATAATAATAA 1302
QY 121 aaagtgtctcgaggagaccttttggcttaaaaagaataataa 180
DB 1303 TAAAAATTAATAATAATAATAATAATAATAATAATAATAATA 1362
QY 181 aaaaagttaataattattattattattattattattattattatt 240
DB 1363 AATTAATAATAATAATAATAATAATAATAATAATAATAATAA 1422
QY 241 taaaaaaatttcaggagggaataataaataataattattcaagtta 300
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RESULT 3

US-08-998-416-186/C

Sequence 186, Application US/08998416
Patent No. 6239264

GENERAL INFORMATION:

APPLICANT: Philippsen, Peter
APPLICANT: Pohlmann, Rainer
APPLICANT: Steiner, Sabine
APPLICANT: Mohr, Christine
APPLICANT: Wendland, Jürgen
APPLICANT: Knechtle, Philipp
TITLE OF INVENTION: GENOMIC DNA SEQUENCES OF ASHBYA GOSSYPII
TITLE OF INVENTION: AND USES THEREOF
NUMBER OF SEQUENCES: 1152
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 6239264artis Corporation
STREET: 3054 Cornwallis Road
CITY: Research Triangle Park
STATE: No. 6239264th Carolina
COUNTRY: USA
ZIP: 27709

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/998,416
FILING DATE: 24-DEC-1997
CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: CH 0016/97
FILING DATE: 31-DEC-1996
ATTORNEY/AGENT INFORMATION:
NAME: Meigs, J. Timothy
REGISTRATION NUMBER: 38,241
REFERENCE/DOCKET NUMBER: PF/5-30306/A/CGC1976
TELECOMMUNICATION INFORMATION:
TELEPHONE: 919-541-8587
TELEFAX: 919-541-8689

INFORMATION FOR SEQ ID NO: 186:

SEQUENCE CHARACTERISTICS:
LENGTH: 615 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: PAG1074RP
US-08-998-416-186

Query Match 18.9%; Score 61.8; DB 4; Length 615;

Best Local Similarity 52.1%; Pred. No. 0.01;
Matches 138; Conservative 0; Mismatches 127; Indels 0; Gaps 0;

QY 62 atataatcatatagctgaataattataatcatatagtaagtaataa 121
DB 596 ATTAAATTAATTAATAATAATAATAATAATAATAATAATAATA 537
QY 122 aagtgtctcgaggagaccttttggcttaaaaagaataataa 181
DB 536 ATTAATAATAATAATAATAATAATAATAATAATAATAATAA 477
QY 182 aaagttaataattattattattattattattattattattatt 241
|||||

TYPE: NUCLEIC ACID

APPLICANT: Philippsen, Peter
 APPLICANT: Pohlmann, Rainer
 APPLICANT: Steiner, Sabine
 APPLICANT: Mohr, Christine
 APPLICANT: Wendland, Jürgen
 APPLICANT: Knechte, Philipp
 APPLICANT: Redischung, Corinne
 TITLE OF INVENTION: GENOMIC DNA SEQUENCES OF ASHBYA GOSSTYPII
 TITLE OF INVENTION: AND USES THEREOF
 NUMBER OF SEQUENCES: 1152
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: No. 6239264artis Corporation
 STREET: 3054 Cornwallis Road
 City: Research Triangle Park
 STATE: No. 6239264th Carolina
 COUNTRY: USA
 ZIP: 27709
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/998,416
 FILING DATE: 24-DEC-1997
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: CH 0016/97

CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.4 Mb storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: Patent in Release #1.0, Version #1.25 (EPO)
CURRENT APPLICATION NUMBER: US/08/617,860B
FILING DATE: 01-MAR-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/EP94/02950

1 PATENT NO.: 3563607
 2
 3 GENERAL INFORMATION:
 4
 5 APPLICANT: Delcuve, Genevieve
 6 APPLICANT: Awang, Gregor
 7 TITLE OF INVENTION: Recombinant DNA Molecules and Expression
 8 TITLE OF INVENTION: Vectors for Tissue Plasminogen Activator
 9 NUMBER OF SEQUENCES: 39
 10
 11 CORRESPONDENCE ADDRESS:
 12 ADDRESSEE: BERESKIN & PARR
 13 STREET: 40 King Street West
 14 CITY: Toronto
 15 STATE: Ontario
 16
 17 COUNTRY: Canada
 18 ZIP: M5H 3Y2
 19
 20 COMPUTER READABLE FORM:
 21 MEDIUM TYPE: Floppy disk
 22 COMPUTER: IBM PC compatible
 23 OPERATING SYSTEM: PC-DOS/MS-DOS
 24 SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/883,795A
FILING DATE: 27-JUN-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Graeville, Michelle
REGISTRATION NUMBER: 40,261
REFERENCE/DOCKET NUMBER: 7841-062
TELECOMMUNICATION INFORMATION:
TELEPHONE: (416) 364-7311
TELEFAX: (416) 361-1398
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 665 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
ORIGINAL SOURCE:
ORGANISM: Homo sapiens
IMMEDIATE SOURCE:
CLONE: Rh 32
US-08-883-795A-36

Query Match 17.0%; Score 55.6; DB 2; Length 665;
Best Local Similarity 50.3%; Pred. No. 0.096;
Matches 163; Conservative 0; Mismatches 159; Indels 4; Gaps 1;

QY 1 attggagatcacttaaatcttagcagaagaatgtttaacgaataaagaataaataa 60
DB 34 ATATTTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 93
QY 61 gatataatataatagctgaataattataatataatgaatgaatgaatgaatga 120
DB 94 AATATTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 153
QY 121 aaagtgcttcggggggaacattttgtttcaaaaggaataataaataattgat 180
DB 154 AATATTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 213
QY 181 aaagtgtaataataatttttttttttttttttttttttttttttttttttt 236
DB 214 AATATTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 273
QY 237 atcgtaaaaaaattcaggggggaataataaagaaaaattttcaagttactgt 296
DB 274 AATGTTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 333
QY 297 aattttatgtttcagtttttttttttttttttttttttttttttttttttt 324
DB 334 AATGTTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 361

RESULT 12
US-08-446-855A-1/c
Sequence 1, Application US/08446855A
Patent No. 5849573
GENERAL INFORMATION:
APPLICANT: Stewart, Thomas S
APPLICANT: Flores, Maria V
APPLICANT: O'Sullivan, William J
TITLE OF INVENTION: Nucleotide sequence encoding carbamoyl
NUMBER OF SEQUENCES: 2
CORRESPONDENCE ADDRESS:
ADDRESSEE: Nixon & Vanderhye PC
STREET: 1100 No. 5849573th Glebe Road, 8th Floor
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22201-4714
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.24
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/446,855A
FILING DATE: 06-JUL-1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mitchard, Leonard C
REGISTRATION NUMBER: 29,009
REFERENCE/DOCKET NUMBER: 47-80
TELECOMMUNICATION INFORMATION:
TELEPHONE: 703-816-4000
TELEFAX: 703-816-4100
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 8920 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: genomic
US-08-446-855A-1

Query Match 16.8%; Score 54.8; DB 2; Length 8920;
Best Local Similarity 53.3%; Pred. No. 0.099;
Matches 163; Conservative 0; Mismatches 137; Indels 6; Gaps 2;

QY 28 aagaatgtttaaatgaataaagaataaagaataaataatataatataatagctga 87
DB 8783 AAAAATTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 8724
QY 88 tataatataatagatagtttaataataaagaatgttcctcggggggaacattttg 147
DB 8723 ATTAATAATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 8664
QY 148 tttaaaaa-----ggaataataaatttgataaagaatgttaataataatttt 203
DB 8663 AATAAATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 8604
QY 204 atttaattgtttaaa--attgataataatgttaaaaaaaatttcaagggggga 261
DB 8603 TGATCATTTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 8544
QY 262 atataaatgaaaaaaattcaagtttcaagtttcaagtttcaagtttcaagttt 321
DB 8543 ATATTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 8484
QY 322 attgtt 327
DB 8483 ACTTTT 8478

RESULT 13
US-09-150-741-1/c
Sequence 1, Application US/09150741
Patent No. 6183996
GENERAL INFORMATION:
APPLICANT: Stewart et al.
TITLE OF INVENTION: Nucleotide Sequence Encoding Carbamoyl Phosphate
FILE REFERENCE:
CURRENT APPLICATION NUMBER: US/09/150,741
CURRENT FILING DATE: 1998-09-10
EARLIER APPLICATION NUMBER: PL6380
EARLIER FILING DATE: 1992-12-16
EARLIER APPLICATION NUMBER: AU93/00617
EARLIER FILING DATE: 1993-12-02
EARLIER APPLICATION NUMBER: 08/446,855
EARLIER FILING DATE: 1995-07-06
NUMBER OF SEQ ID NOS: 15

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[illegible]

PS Example 6; Page 58; 101pp; English.
XX
CC Compositions containing neutralising antitoxin against one or more E.
CC coli verotoxin (VT) can be used to treat intoxicated adults and
CC children with enteric bacterial infections. They may also be used as
CC prophylactics e.g. as a vaccine, against diarrhoeal disease or the
CC development of extra-intestinal complications of E.coli infection,
CC especially haemolytic uraemic syndrome. The antitoxin can also be
CC used to detect E. coli VT in a sample. The VT is recombinant,
CC preferably a fusion protein containing a non-VT protein sequence and
CC part of the E.coli VT1 or VT2 sequence. Two primers (AA12655,
CC AA12656) were used to amplify the verotoxin VT-1 A subunit coding
CC sequence and add a histidine tag coding sequence to the subunit
CC sequence. Two primers (AA12655, AA12658) were used to amplify the
CC verotoxin VT-1 A and B subunits and add a histidine tag coding
CC sequence to the subunit sequences.
SQ Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.9%; Score 16; DB 17; Length 29;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 189 aaaaataattatttta 204
Db 9 aaaaataattatttta 24

RESULT 2
ID AAA51200 standard; DNA; 29 BP.
XX
AC AAA51200;
XX
DT 26-SEP-2000 (first entry)
XX
DE N-terminal primer for E. coli verotoxin 1 subunit A gene.
XX
KM VT-1; verotoxin; antitoxin therapy; fusion protein; affinity tag; food;
KM recombinant production; screening; dairy; anti-bacterial; vaccine;
KM primer; polystyrene; ss.
XX
OS Escherichia coli.
OS Synthetic.
XX
PN US6080400-A.
XX
PD 27-JUN-2000.
XX
PF 13-MAR-1997; 97US-0816977.
XX
PR 24-MAR-1995; 95US-0410058.
XX
PA (OPHI-) OPHIDIAN PHARM INC.
XX
PI Williams JA, Byrne LM;
XX
DR WPI; 2000-451195/39.
XX
XX Bacterial cell for recombinantly expressing bacterial toxins in large
PT quantities useful for immunization and treatment of bacterial
PT infections, comprises expression vector encoding bacterial toxin
XX
PS Example 6; Column 83; 83pp; English.
XX
CC E. coli verotoxin (VT) type 1 and 2 subunits A and B were cloned into
CC pET-23b, designed to allow expression of the native proteins containing
CC C-terminal polystyrene tags. The VT-1 and VT-2 genes were engineered
CC to convert the signal sequence methionine codon into a NdeI site to
CC allow cloning of the amplified genes into the vector without addition of
CC vector-encoded amino acids. The C-terminal primers comprises the
CC C-terminal 7 codons of each gene fused to the sequence CTCGAGCC, in order

CC to add the polystyrene tag. The primers delete the native stop codons,
CC and when cloned into pET-23 add a C-terminal extension of Leu-Glu-(His)₆.
CC VT B chains are small proteins (approximately 8 kDa), so use of a small
CC affinity tag was preferred (i.e. polystyrene). A polystyrene affinity
CC tag facilitates single step affinity purification of subunits from
CC periplasmic extracts. However, due to poor recovery of his-tagged VT-1 A
CC and VT-2 A chains, expression of maltose binding protein (MBP) fused
CC subunits was undertaken. Due to the toxicity of the VT-2 B subunit,
CC strict uninduced promoter control is necessary to permit cell viability.
CC Bacterial host cells expressing a recombinant expression vector encoding
CC a polystyrene affinity tag and a portion of the VT-2 B chain are
CC claimed. The vector is chosen from pET24hisVT2BL+, pET24hisVT2BL- and
CC pET24VT2B, where "L+" indicates that the vector encodes the preprotein
CC form of the protein and "L-" indicates that the vector encodes the mature
CC form of the protein. The bacterial cell is capable of expressing large
CC quantities (40 mg/l) of VT-2B. The toxins are useful for immunizing
CC non-mammals and for detecting bacterial toxins in environmental samples
CC including soil, water, industrial samples, biological samples and samples
CC obtained from food and dairy processing instruments.
SQ Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.9%; Score 16; DB 21; Length 29;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 189 aaaaataattatttta 204
Db 9 aaaaataattatttta 24

RESULT 3
ID AA227689 standard; DNA; 32 BP.
XX
AC AA227689;
XX
DT 22-DEC-1999 (first entry)
XX
DE PCR primer for Verotoxin gene.
XX
KM Verotoxin; VT1; VT2; detection; PCR primer; ss.
XX
OS Synthetic.
OS Escherichia coli.
XX
PN JP11243996-A.
XX
PD 14-SEP-1999.
XX
PF 27-FEB-1998; 98JP-0047677.
XX
PR 27-FEB-1998; 98JP-0047677.
XX
PA (TOYM) TOYOMO KK.
XX
DR WPI; 1999-603716/52.
XX
XX An oligonucleotide for amplification of verotoxin - useful in the
PT detection of inactivated verotoxin gene by transfer of a foreign DNA
PT fragment
XX
PS Claim 11; Page 9; 10pp; Japanese.
XX
CC This sequence represents a PCR primer of the invention. The primer is
CC used for amplification of the E. coli verotoxin (VT) gene. The
CC oligonucleotide is useful for detection of inactivated VT gene by
CC transfer of a foreign DNA fragment. Simple, rapid and specific
CC amplification of VT gene from environmental factors is achieved using the
CC oligonucleotide of the invention.
XX
SQ Sequence 32 BP; 12 A; 2 C; 4 G; 14 T; 0 other;

Query Match 4.9%; Score 16; DB 20; Length 32;
 Best Local Similarity 100.0%; Pred. No. 1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
 |||||
 Db 4 aaataattatttta 19

RESULT 4

AA97603
 ID AAT97603 standard; DNA; 36 BP.

XX AAT97603;

XX 30-APR-1998 (first entry)

XX Shigella dysenteriae delta-stxA allele PCR primer 13.

XX Delta-virg allele; delta-guab-A allele; PCR: amplification; primer;
 XX delta-stxA allele; shigellosis; vaccine; ss.

XX Synthetic.

XX Shigella dysenteriae.

XX W09737685-A1.

XX 16-OCT-1997.

XX 09-APR-1997; 97WO-US05954.

XX 09-APR-1996; 96US-0629600.

XX (UYMA-) UNIV MARYLAND BALTIMORE.

XX Levine MM, Noriega FR;

XX WPI; 1997-512417/47.

XX Shigella mutants with mutation in guab-A - used in vaccines against
 XX Shigellosis

XX Example 6; Page 57; 94pp; English.

XX This is a PCR primer used in the amplification of the Shigella
 XX dysenteriae 1 delta-stxA allele. The delta-stxA allele was integrated
 XX into delta-guab-A of delta-guab-A, delta-virg S. dysenteriae 1, which
 XX inactivated the shiga toxin of this strain. The mutant can be used in
 XX the preparation of vaccines such as, a live vector vaccine comprising
 XX a Shigella mutant, (which encodes and expresses a foreign antigen, and
 XX a pharmaceutically acceptable carrier) or a DNA mediated vaccine
 XX comprising the Shigella mutant (which also contains a plasmid which
 XX encodes and expresses a foreign antigen in a eukaryotic cell). The
 XX vaccines can be used against Shigellosis.

XX Sequence 36 BP; 11 A; 3 C; 10 G; 12 T; 0 other;

Query Match 4.9%; Score 16; DB 18; Length 36;
 Best Local Similarity 100.0%; Pred. No. 1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
 |||||
 Db 14 aaataattatttta 29

RESULT 5

AA90775

ID AAV90775 standard; DNA; 39 BP.
 XX

AC AAV90775;

XX 18-FEB-1999 (first entry)

XX Primer Y104F.

XX Antigen; immunogenic cluster family; vaccine; gastritis; diagnosis;
 XX peptic ulcer; gastric adenocarcinoma; gastric lymphoma; primer; ss.

XX Synthetic.

XX W09849314-A2.

XX 05-NOV-1998.

XX 27-APR-1998; 98WO-US08487.

XX 14-OCT-1997; 97US-0061958.

XX 25-APR-1997; 97US-0045107.

XX (GENE-) GENELABS TECHNOLOGIES INC.

XX Chow TP, Fry KE, Lim MY, McAttee CP;

XX WPI; 1999-009433/01.

XX New Helicobacter pylori antigens and related nucleic acid sequences
 XX - useful in serological diagnosis and protective vaccines, providing
 XX long-lasting immune response
 XX Claim Disclosure; Page 194; 402pp; English.

XX The specification, which describes Helicobacter pylori antigenic
 XX proteins that are characterised by immunoreactivity with
 XX H. pylori-positive antisera. The specification also describes 69
 XX previously unrecognised immunogenic cluster families. H. pylori
 XX antigens are used to detect H. pylori-specific antibodies, for
 XX diagnosing infection or to confirm eradication of infection, and
 XX in vaccines to protect against H. pylori infection and related
 XX diseases (gastritis, peptic ulcer, gastric adenocarcinoma/lymphoma).
 XX The present primer is used in the course of the invention.

XX Sequence 39 BP; 15 A; 8 C; 7 G; 9 T; 0 other;

Query Match 4.9%; Score 16; DB 20; Length 39;
 Best Local Similarity 100.0%; Pred. No. 1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 8 atacttaatttcg 23
 |||||
 Db 23 atacttaatttcg 38

RESULT 6

AAZ68379
 ID AAZ68379 standard; DNA; 47 BP.

XX AAZ68379;

XX 10-SEP-2001 (first entry)

XX Human map-related biallelic marker SEQ ID NO:2726.

XX Human genome; biallelic marker; high density disequilibrium map;
 XX genomic map; haplotype; phenotype; polymorphic base; genotyping;
 XX haplotyping; hybridisation; identification; characterisation;
 XX diagnosis; single nucleotide polymorphism; SNP; ds.

XX Homo sapiens.

XX Key Location/Qualifiers
 XX variation replace(24,A)
 FT

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FT      /*tag= a
XX      /standard_name= "single nucleotide polymorphism"
XX      W09954500-A2.
XX      PD
XX      PD      28-OCT-1999.
XX      PF      21-APR-1999; 99WO-IB00822.
XX      PR      21-APR-1998; 98US-0082614.
XX      PR      23-NOV-1998; 98US-0109732.
XX      PA      (GEST ) GENSET.
XX      PI      Cohen D, Blumenfeld M, Chumakov I;
XX      DR      WPI: 2000-013267/01.
XX      PT      Novel biallelic markers used to construct a high density disequilibrium
XX      PT      map of the human genome -
XX      PS      Claim 3; Page 812; 2745pp; English.
XX      CC      AA265654 to AA269578 represent human biallelic markers from the present
XX      CC      invention, which contain a polymorphic base at position 24 of their
XX      CC      nucleotide sequences. AA269579 to AA277440 represent amplification
XX      CC      primers for the biallelic markers. The biallelic markers of the
XX      CC      invention have a variety of uses: they can be used for high density
XX      CC      mapping of the human genome, and in complex association studies and
XX      CC      haplotyping studies which are useful in determining the genetic basis
XX      CC      for disease states. Compositions and methods of the invention can also
XX      CC      be useful for the identification of the targets for the development of
XX      CC      pharmaceutical agents and diagnostic methods, as well as the
XX      CC      characterisation of the differential efficacious responses to and side
XX      CC      effects from pharmaceutical agents acting on a disease as well as other
XX      CC      treatment.
XX      CC      N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
XX      CC      and 3367, are not actually given a sequence in the Sequence Listing
XX      CC      from the present invention.
XX      SQ      Sequence 47 BP; 24 A; 5 C; 6 G; 12 T; 0 other;

Query Match      4.9%; Score 16; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 9.8e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      147 gtcttaaaaggagaaa 162
        |||||||||||||||
DB      13 gttttaaaaggagaaa 28

RESULT 7
ID      AAA66183 standard; DNA; 20 BP.
XX      AC      AAA66183;
XX      DT      09-OCT-2000 (first entry)
XX      DE      Dog genomic marker oligonucleotide sequence SEQ ID NO:45.
XX      KW      Dog; genome; genomic marker; radiation hybrid map; identification;
XX      KW      chromosome location; gene marker; polymorphic microsatellite marker;
XX      KW      phenotype; behaviour; pedigree; ss.
XX      OS      Canis familiaris.
XX      PN      WO200029615-A2.
XX      PD      25-MAY-2000.
XX      PF      15-NOV-1999; 99WO-IB01907.

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XX      13-NOV-1998; 98US-0108193.
XX      PR      (CNRS ) CNRS CENT NAT RECH SCI.
XX      PI      Gallibert F, Andre C;
XX      DR      WPI: 2000-387821/33.
XX      PT      New radiation hybrid map of the dog, Canine familiaris, genome, useful
XX      PT      for e.g. identifying genes implicated in phenotypic and behavioral
XX      PT      traits or in genetic diseases and for studying dog pedigrees -
XX      PS      Claim 1; Page 55; 87pp; English.
XX      CC      The present invention describes a radiation hybrid map of the dog
XX      CC      (Canine familiaris) genome comprising the genome location of a marker
XX      CC      selected from AA66139 to AA66942. The radiation hybrid map is useful
XX      CC      for identifying and localising dog genes, since it covers approximately
XX      CC      80 % of the dog genome and provides a dense map integrating different
XX      CC      types (i.e. Type I and Type II) of markers. The map and the dog genome
XX      CC      markers (or complementary sequences) are especially useful to identify
XX      CC      genes responsible for phenotypic and behavioural traits in dogs, to
XX      CC      identify morbid genes, to analyse diseases and identify implicated genes
XX      CC      in such diseases and their alleles, and to study dog pedigrees. They
XX      CC      may also be useful for isolating corresponding human gene sequences
XX      CC      e.g. genes involved in genetic diseases.
XX      SQ      Sequence 20 BP; 14 A; 1 C; 2 G; 3 T; 0 other;

Query Match      4.6%; Score 15; DB 21; Length 20;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      261 aatataaatgagaaa 275
        |||||||||||||||
DB      1 aatataaatgagaaa 15

RESULT 8
ID      AAH46804 standard; DNA; 24 BP.
XX      AC      AAH46804;
XX      DT      19-SEP-2001 (first entry)
XX      DE      Human high motility group protein family 11 cDNA PCR primer #2.
XX      DE      Human; high motility group protein family 11; cancer; haemopathy;
XX      KW      HIV infection; immune disease; inflammation; gene therapy;
XX      KW      PCR primer; ss.
XX      OS      Homo sapiens.
XX      PN      WO200147967-A1.
XX      PD      05-JUL-2001.
XX      PF      18-DEC-2000; 2000WO-CN00595.
XX      PR      23-DEC-1999; 99CN-0125721.
XX      PA      (UYFU-) UNIV FUDAN.
XX      PA      (SHAN-) SHANGHAI BIO DOOR GENE TECHNOLOGY LTD.
XX      PI      Mao Y, Xie Y;
XX      DR      WPI: 2001-418226/44.
XX      PT      High motility group protein family 11 and encoded polynucleotide,
XX      PT      applicable in diagnosis and treatment of cancer, haemopathy, HIV

```



```

PT infection, immunological diseases and various inflammation
XX
PS Example 3, Page 17; 39pp; Chinese.
XX
CC The present invention provides the protein and coding sequences of the
CC human high motility group protein family 11. The sequences are useful in
CC the treatment of cancer, haemopathy, HIV infection, immune diseases and
CC inflammation. The present sequence is a PCR primer for the coding
CC sequence of the invention.
XX
SQ Sequence 24 BP; 8 A; 3 C; 0 G; 13 T; 0 other;

Query Match
Best Local Similarity 100.0%; Pred. No. 2.6e+03; Length 24;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 193 taattattttattt 207
   |||||||
Db 2 taattattttattt 16

RESULT 9
AAT39439
ID AAT39439 standard; RNA; 27 BP.
XX
AC AAT39439;
XX
DT 13-NOV-1996 (first entry)
XX
DE Hel-N2 selected sequence, e-13.
XX
KW Human: neuron-specific protein; Hel-N1; 3'-UTR instability sequence;
KW paraneoplastic sensory neuropathy; oncoprotein; lymphokine; rat; elav;
KW RNA recognition motif; RRM; Drosophila; cellular growth; localisation;
KW instability; translatibility; neurons; autoimmune protein; PE; PCD; PSN;
KW central nervous system; cancer; paraneoplastic cerebellar degeneration;
KW paraneoplastic encephalomyelitis; RNP-1 octamer sequence;
KW human elav-like neuronal protein-1; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_RNA 15..23 /*tag= a
FT /note= "Consensus sequence"
XX
PN US5525495-A.
XX
PD 11-JUN-1996.
XX
PE 11-MAY-1992; 92US-0881075.
XX
PR 15-SEP-1993; 93US-0120827.
PR 11-MAY-1992; 92US-0881075.
XX
PA (UYDU-) UNIV DUKE.
XX
PI Gao F, Keene JD, Levine T;
XX
DR WPI: 1996-286398/29.
XX
PT Prodn of cDNA library for related proteins - by screening total cell
XX mRNA with RNA-binding protein Hel-N1 or Hel-N2
XX
PS Disclosure; Fig 11; 66pp; English.
XX
CC The sequences given in AAT39427-46 represent random oligonucleotides
CC which were isolated due to their ability to bind to the human neuron-
CC specific protein, Hel-N2. These sequences contain short stretches
CC of uridylylate residues interspersed with other residues. These U-
CC rich regions share homology with the 3'-UTR instability sequences
CC that are found in mRNA's. Instability sequences are target elements

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CC which reside in the 3'-non-coding regions of mRNA's which encode
CC oncoproteins and lymphokines. Hel-N2 is a deleted form of Hel-N1
CC in which the residues 239-251 have been deleted. This protein is
CC expressed in medulloblastoma tumour cells and is not found in whole
CC human brain. A small amount of Hel-N2 is also found in fetal brain,
CC which may indicate a correlation with rapid growth. Hel-N1 cDNA was
CC isolated by probing for rat and human elav counterparts using degenerate
CC primers designed to simulate the RNP-1 octamer sequence present in two
CC of the three RNA recognition motifs (RRM's) of Drosophila elav. Hel-N1
CC was found to contain 3 RRM's, where the third one (see also AAM00244) is
CC sufficient for mRNA 3'-UTR binding activity. Full length Hel-N1, when
CC transfected into a cell, causes cellular growth to cease, however, if
CC just the third binding domain is transfected into cells, the cells
CC undergo rapid growth. Hel-N1 binds as a multimer along the mRNA,
CC presumably enhancing its localisation, instability and/or regulating it
CC translatibility and/or deadenylation. This protein may be
CC responsible for the growth cessation of neurons. Hel-N1 is an autoimmune
CC protein in certain patients who show central nervous system
CC manifestations of cancer called paraneoplastic cerebellar degeneration
CC (PCD), paraneoplastic encephalomyelitis (PE) or paraneoplastic sensory
CC neuropathy (PSN).
XX
SQ Sequence 27 BP; 7 A; 2 C; 2 G; 16 U; 0 other;

Query Match
Best Local Similarity 33.3%; Pred. No. 2.6e+03; Length 27;
Matches 5; Conservative 10; Mismatches 0; Indels 0; Gaps 0;

OY 197 tattttattttaa 211
   :|:::|:|:::|
Db 12 uauuuuuuuuuuaa 26

RESULT 10
AAV37457
ID AAV37457 standard; RNA; 27 BP.
XX
AC AAV37457;
XX
DT 07-SEP-1998 (first entry)
XX
DE Human Hel-N2 selected RNA sequence e-13.
XX
KW Growth regulatory protein; Hel-N2; oncogene; cytokine; lymphokine;
KW chromosome mapping; human; functionally related protein; ss.
XX
OS Homo sapiens.
XX
PN US5773246-A.
XX
PD 30-JUN-1998.
XX
PE 07-JUN-1995; 95US-0478675.
XX
PR 15-SEP-1993; 93US-0120827.
PR 11-MAY-1992; 92US-0881075.
PR 07-JUN-1995; 95US-0478675.
XX
PA (GAOF/) GAO F,
PA (KEEN/) KEENE J D,
PA (LEVI/) LEVINE T.
XX
PI Gao F, Keene JD, Levine T;
XX
DR WPI: 1998-387003/33.
XX
PT Use of proteins which bind RNA - for obtaining a cDNA library
XX containing members encoding structurally or functionally related
XX proteins from total cell mRNA.
XX
PS Disclosure; Fig 11; 67pp; English.
XX

```

CC This represents a RNA sequence selected from a randomised RNA library
CC by binding to human Hel-N2 under high stringency conditions. The
CC invention provides a method for obtaining a cDNA library having members
CC encoding a group of structurally or functionally related proteins from
CC total cell mRNA. The method comprises binding RNA representing total cell
CC mRNA to a protein that binds RNA and has specific binding to untranslated
CC regions of a subset of the total cell mRNA, where the protein that binds
CC RNA has been purified to remove any other protein that binds RNA. The
CC resulting bound products are separated and a cDNA library is prepared.
CC The protein that binds RNA may be Hel-N1, Hel-N2, Carg, DT-7, K1, K2,
CC K3, Hud, HNC, elav, rbp9, e1f4b, sxl, tra-2, AUBF, AUF, ASF/SF2, U2AF,
CC SC35, or hnRNP proteins. The cDNAs obtained can be used for chromosome
CC mapping and genome sequencing of the structurally or functionally related
CC genes encoding growth regulatory proteins, proto-oncogenes, cytokines,
CC lymphokines, or anti-oncogene proteins. They can also be used in
CC diagnostic methods for determining imbalances in such genes and the
CC efficacy of various treatments to correct any imbalance.

XX Sequence 27 BP; 7 A; 2 C; 2 G; 16 U; 0 other;

Query Match 4.6%; Score 15; DB 19; Length 27;
Best Local Similarity 33.3%; Pred. No. 2.6e+03;
Matches 5; Conservative 10; Mismatches 0; Indels 0; Gaps 0;

OY 197 tattttattttaa 211
:|:::|:|:|:|:|

DB 12 uuuuuuuuuuuuaa 26

RESULT 11

AA043975
ID AA043975 standard; DNA; 30 BP.

XX AA043975;

DF 28-OCT-1993 (first entry)

DE Triple helix forming oligonucleotide III.

KM Purine; pyrimidine; tracts; intramolecular triplex; therapeutic;
KM diagnostic; control; gene expression; mRNA synthesis suppression;
KM ss.

XX Synthetic.

OS WO9312230-A.

PM 24-JUN-1993.

PD 11-DEC-1992; 92MO-US10792.

PF 13-DEC-1991; 91US-0808452.

PR 21-JAN-1992; 92US-0826934.

XX (STRI) SRI INT.

XX Jayasena SD, Johnston BH;

PI WPI; 1993-214172/26.

DR New oligo:nucleotide(s) forming triple helix with target nucleic
XX acid - contain purine and pyrimidine tracts in specific
XX orientations, useful therapeutically or diagnostically e.g. for
XX inactivating HIV RNA, etc.

XX Disclosure: Page 48; 101pp; English.

XX The sequence is that of an oligonucleotide, III, which is able to form
CC a triple helix with a duplex nucleic acid (dsDNA) contg. a target
CC sequence which comprises at least one pyrimidine tract, and at least
CC one adjacent purine tract. It is useful for therapeutic or
CC diagnostic control of gene expression, e.g. suppression of mRNA

CC synthesis from a target gene. A specified application is targeting
CC of RNA in the HIV-1 genome. When appropriately labelled it may also
CC be used as a probe. Attachment of cleavage agents caused permanent
CC inactivation of the target by site-specific cleavage.

XX Sequence 30 BP; 12 A; 0 C; 5 G; 13 T; 0 other;

Query Match 4.6%; Score 15; DB 14; Length 30;
Best Local Similarity 100.0%; Pred. No. 2.6e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 212 ttgtgttaaaatttg 226
|||||

DB 5 ttgtgttaaaatttg 19

RESULT 12

AA025576
ID AA025576 standard; cDNA to mRNA; 48 BP.

XX AA025576;

DF 15-NOV-1996 (first entry)

DE Human gene signature HMWGS07754.

KM Gene signature; messenger RNA; mRNA; relative abundance; frequency;
KM human; cloning; mapping; non-biased library; diagnosis; detection;
KM cell typing; abnormal cell function; ss.

XX Homo sapiens.

OS WO9514772-A1.

PM 01-JUN-1995.

PD 11-NOV-1994; 94MO-JP01916.

PF 12-NOV-1993; 93JP-0355504.

PR (MATS/) MATSUBARA K.

PA (OKUB/) OKUBO K.

XX Matsubara K, Okubo K;

PI WPI; 1995-206931/27.

DR Identifying gene signatures in 3'-directed human cDNA library - e.g.
XX for diagnosis of abnormal cell function, by preparing cDNA that
XX reflects relative abundance of corresp. mRNA in specific human
XX tissues

PS Claim 1; Page 1862; 2245pp; Japanese.

XX A single-stranded DNA (or its complementary strand or the corresp.
XX double-stranded DNA) which comprises one of the 7837 "GS" sequences
XX given in AA019001-1726837 and which is able to hybridise to part of
XX human genomic DNA, cDNA or mRNA is claimed. The GS (gene signature)
XX sequences were obtained from 3'-directed cDNA libraries prepared
XX from various human tissues; synthesis of cDNA was initiated from the
XX 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
XX untranslated sequence is unique to a particular mRNA species, almost
XX all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
XX is constructed so as to reflect accurately the relative abundance of
XX different mRNAs in the particular tissue from which it was derived.
XX The appearance frequency of a given GS in a cDNA library can be
XX determined (esp. using primers and probes derived from the GS
XX sequences) as a means of diagnosing abnormal cell function or for
XX recognising different cell types.

XX Sequence 48 BP; 26 A; 5 C; 4 G; 13 T; 0 other;

Query Match 4.6%; Score 15; DB 16; Length 48;
Best Local Similarity 100.0%; Pred. No. 2.5e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 159 aaatataataaa 173
|||||
Db 17 aaatataataaa 31

RESULT 13
AAQ20161
ID AAQ20161 standard; DNA; 18 BP.
XX
AC AAQ20161;
XX
DT 01-APR-1992 (first entry)
XX
DE Cross-linking oligomer 724 to target Herpes Simplex Virus I.
XX
KW deoxyribonucleic acid; major groove; HSV;
XX Inverted polarity region; covalent cross-linking group; ss.
XX Synthetic.
XX
FH Key
FT modified_base 1 location/Qualifiers
FT /*tag= a
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 2
FT /*tag= b
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 3
FT /*tag= c
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 4
FT /*tag= d
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 5
FT /*tag= e
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 7
FT /*tag= f
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 9
FT /*tag= g
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 10
FT /*tag= h
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 11
FT /*tag= i
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT misc_feature 12..18
FT /*tag= j
FT /label= inverted_polarity_region
FT /note= "see comments"
FT 13
FT /*tag= k
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 14
FT /*tag= l
FT /mod_base= OTHER

FT modified_base 15 /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT /*tag= m
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 17
FT /*tag= n
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT
FT WO9118997-A.
XX 12-DEC-1991.
XX
XX 24-MAY-1991; 91WO-1003680.
XX
XX 14-JAN-1991; 91US-0640654.
XX 25-MAY-1990; 90US-0529346.
XX
XX (GILE-) GILEAD SCIE INC.
XX
XX Matteucci MD, Krawczyk S;
XX WPI; 1992-007480/01.
XX
XX
XX New sequence-specific non-photo-activated crosslinking agents -
XX bind to the major groove of duplex DNA and are esp. useful for
XX treating latent infections e.g. HIV
XX
XX Example 4; Page 29; 42pp; English.
XX
XX This oligomer contains an inverted polarity region formed from an
XX o'-xyloso dimer synthon. Residues 11 and 12 are linked via an
XX o'-xyloso group (i.e. nucleotides that have xylose sugar linked via
XX the o'-xylene ring). The sequence is designed to target the Herpes
XX Simplex virus I beginning at nucleotide 10996 and to covalently
XX cross-link to it. See also AAQ20151-Q20160.
XX
XX Sequence 18 BP; 12 A; 1 C; 0 G; 5 T; 0 other;
XX SQ

Query Match 4.3%; Score 14; DB 13; Length 18;
Best Local Similarity 100.0%; Pred. No. 6.9e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 159 aaatataataaa 172
|||||
Db 2 aaatataataaa 15

RESULT 14
AAQ20160
ID AAQ20160 standard; DNA; 18 BP.
XX
AC AAQ20160;
XX
DT 01-APR-1992 (first entry)
XX
DE Cross-linking oligomer 723 to target Herpes Simplex Virus I.
XX
KW deoxyribonucleic acid; major groove; HSV;
XX Inverted polarity region; covalent cross-linking group; ss.
XX Synthetic.
XX
FH Key
FT modified_base 1 location/Qualifiers
FT /*tag= a
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT modified_base 2
FT /*tag= b
FT /mod_base= OTHER

FT		/note= "N-methyl-8-oxo-2'-deoxyadenine"
ET	modified_base	3
FT		/*tag= c
FT	/mod_base= OTHER	
ET	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	4
FT		/*tag= d
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	5
FT		/*tag= e
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	7
FT		/*tag= f
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	9
FT		/*tag= g
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	10
FT		/*tag= h
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	11
FT		/*tag= i
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	misc_feature	12..18
FT		/*tag= j
FT	/label= Inverted_polarity_region	
FT	/note= "see comments"	
FT	modified_base	13
FT		/*tag= k
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	14
FT		/*tag= l
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	15
FT		/*tag= m
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
FT	modified_base	17
FT		/*tag= n
FT	/mod_base= OTHER	
FT	/note= "N-methyl-8-oxo-2'-deoxyadenine"	
XX		
PN	W09118997-A.	
PD	12-DEC-1991.	
XX		
PP	24-MAY-1991;	91WO-1003680.
XX		
PR	14-JAN-1991;	91US-0640654.
PR	25-MAY-1990;	90US-0529346.
XX		
PA	(GILE-) GILEAD SCIE INC.	
PI		
PI	Matteucci MD, Krawczyk S;	
DR	WPI; 1992-007480/01.	
CC		

This oligomer contains an inverted polarity region formed from o-xylolo dimer synthon. Residues 11 and 12 are linked via an

[illegible]

FT		modified_base	14	/tag= k
PT				/mod_base= OTHER
ET				/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT		modified_base	15	/tag= l
PT				/mod_base= OTHER
ET				/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT		modified_base	17	/tag= m
PT				/mod_base= OTHER
ET				/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT		misc_feature	12..18	/tag= n
PT				/label= inverted.polarity_region
ET				/note= "see comments"
FT		misc_feature	11..12	/tag= o
PT				/note= "O'-xylosio dimer synthon linkage"
XX				
PB		WO9209705-A.		
XX				
PD		11-JUN-1992.		
XX				
PF		25-NOV-1991;	91WO-US08811.	
XX				
PR		23-NOV-1990;	90US-0617907.	
PR		18-JAN-1991;	91US-0643382.	
PR		08-APR-1991;	91US-0683420.	
PR		17-APR-1991;	91US-0686544.	
PR		17-APR-1991;	91US-0686546.	
PR		17-SEP-1991;	91US-0686547.	
PR			91US-0766733.	
PA	(GILEAD)	GILEAD SCI INC.		
PI	Froehner B,	Krawczyk S,	Matteucci MD,	Malligan J;
DR	WPl:	1992-217083/26.		
XX		New oligomers contg. modified bases - which form a triplex with G-C doublet in a DNA duplex, for treating and diagnosing HIV, hepatitis, herpes, malignancy and inflammation		
PS	Claim 12; Page 67; 77pp;	English.		
CC	The synthetic oligomer is capable of forming a triplex at physiological pH with a purine rich target sequence by coupling into the major groove of the duplex. The specific target sequence of this oligomer is a herpes simplex virus I duplex beginning at nucleotide 10996 contg. a purine-rich region concentrated on one chain of the duplex. The oligomer, and others like it are useful in diagnosis and therapy of diseases characterised by specific DNA duplex targets, e.g. respiratory syncytial virus, HIV, hepatitis, herpes, malignant tumours and inflammation. The triple helices form under mild conditions thus assays may be carried out without subjecting the test specimen to harsh conditions. The oligomer contains an inverted polarity region formed from an O'-xyliso diener synthon. The linking gp. is O'-xyliso (nucleotides have the 3' positions of xlyose sugars linked via the O'-ylene ring). Two nucleotides are coupled through a xylen residue to form the dimer synthon. This additional modifications may render the oligmer stable to nuclease activity. The oligomer is able to inhibit gene expression, as verified by in-vitro systems.			
CC	See also AAQ25452-25501 and AAQ30226-448.			
SQ	Sequence 18 BP; 12 A; 1 C; 0 G; 5 T; 0 other;			
Query Match	Best Local Similarity	4.3%; Score 14; DB 13; Length 18; Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		

Qy	159	aaatataataaa	172
Db	2	aaaataataaaa	15
RESULT	16		
ID	AAQ30310	standard; DNA; 18 BP.	
XX	AAQ30310;		
XX	07-DEC-1992	(first entry)	
DE	Oligomer HSV723	for forming triplex with HSV target duplex.	
XX	Herpes simplex virus I; AIDS; modified;	HIV; RSV; HPV; malignant	
KW	hepatitis; inflammation; ss.		
XX	Synthetic..		
OS			
Key	modified_base	Location/Qualifiers	
FH		1 /tag= a	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		2 /tag= b	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		3 /tag= c	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		4 /tag= d	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		5 /tag= e	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		6 /tag= f	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		7 /tag= g	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		8 /tag= h	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		9 /tag= i	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		10 /tag= j	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		11 /tag= k	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		12 /tag= l	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		13 /tag= m	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		14 /tag= n	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		15 /tag= o	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		16 /tag= p	
FT		/mod_base= OTHER	
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FT		18 /tag= r	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		19 /tag= s	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		20 /tag= t	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		21 /tag= u	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		22 /tag= v	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		23 /tag= w	
FT		/mod_base= OTHER	
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FT		24 /tag= x	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		25 /tag= y	
FT		/mod_base= OTHER	
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FT		26 /tag= z	
FT		/mod_base= OTHER	
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FT		27 /tag= AA	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		28 /tag= AB	
FT		/mod_base= OTHER	
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FT		31 /tag= AE	
FT		/mod_base= OTHER	
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FT		32 /tag= AF	
FT		/mod_base= OTHER	
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FT		33 /tag= AG	
FT		/mod_base= OTHER	
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FT		34 /tag= AH	
FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		35 /tag= AI	
FT		/mod_base= OTHER	
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FT		/mod_base= OTHER	
FT		/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT		37 /tag= AK	


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XX 22-MAR-2001.
PD
XX
XX 14-SEP-2000; 2000WO-US25479.
PF
XX
XX 15-SEP-1999; 99US-0398522.
PR
XX
PA (UYJO ) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
XX
XX Issa J;
PI
XX
XX WPI; 2001-244777/25.
DR
XX
PT New nucleic acid molecule for use as a marker for screening cancer,
PT comprises the coding region for a T-type calcium channel and regulatory
PT sequences associated with the channel
XX
XX
XX Claim 21; Page 34; 125pp; English.
XX
XX The present sequence for 5'-bisulfite PCR primer is used to study the
CC methylation state of human RasGAP-related protein (IQGAP2) which
CC maps to chromosome 5q. The methylation state of specific regions
CC within CpG islands associated with a novel T-type calcium channel
CC CACNA1G gene correlate with several cancerous phenotypes involving
CC various tissue and cell types. Since aberrant methylation of normally
CC unmethylated CpG islands is often observed in immortalised and
CC transformed cells, CACNA1G is implicated in cellular proliferative
CC disorders e.g. leukaemia, colorectal, lung, breast and other cancers. The
CC nuclear acid coding for CACNA1G is useful as a marker for screening
CC cancer and age related diseases. A diagnostic kit containing primers
CC (AAS01574-AAS01623) for amplification of a CpG-containing nucleic acid,
CC where the primer hybridises with a target polynucleotide sequence
CC (AAS01627-AAS01676), can be used for detecting aberrant methylation. The
CC CpG island sequences (AAS01677-AAS01692) are selected from genes encoding
CC CACNA1G, apolipoprotein B (APOB), caudal type homeobox transcription
CC factor 2 (CDX2), epidermal growth factor receptor (EGFR), fibrillin-1
CC (FBN1), G protein-coupled receptor 37 (GPR37), heat shock 70kD protein 6
CC (HSP70B), HSPA6), RasGAP-related protein (IQGAP2), Klotho (KL),
CC proteinase-activated receptor 2 (PAR2), paired-like homeodomain
CC transcription factor 2 (PTRX2), patched A and B (PTCHA; PTCHB) and
CC syndecan 1 and 4 (SDC1; SDC4) or a MINT31 sequence.
XX
XX Sequence 22 BP; 4 A; 0 C; 3 G; 15 T; 0 other;
SQ
Query Match 4.38; Score 14; DB 22; Length 22;
Best Local Similarity 100.0%; Pred. No. 6,8e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 196 ttattttatttta 209
Db 2 ttattttatttta 15

```

```

XX OS Homo sapiens.
XX
XX PN W0200119645-A1.
XX
XX PD 22-MAR-2001.
XX
XX PF 14-SEP-2000; 2000WO-US25479.
XX
XX PR 15-SEP-1999; 99US-0398522.
XX
XX PA (UYJO ) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
XX
XX Issa J;
PI
XX
XX WPI; 2001-244777/25.
DR
XX
PT New nucleic acid molecule for use as a marker for screening cancer,
PT comprises the coding region for a T-type calcium channel and regulatory
PT sequences associated with the channel
XX
XX
XX Claim 20; Page 37; 125pp; English.
XX
XX The present sequence for human RasGAP-related protein (IQGAP2)
CC 5'-target sequence (complementary to the 5'-bisulfite PCR primer) is used
CC to study the methylation state of IQGAP2 which maps to chromosome 5q. The
CC methylation state of specific regions within CpG islands associated with
CC a novel T-type calcium channel CACNA1G gene correlate with several
CC cancerous phenotypes involving various tissue and cell types. Since
CC aberrant methylation of normally unmethylated CpG islands is often
CC observed in immortalised and transformed cells, CACNA1G is implicated in
CC cellular proliferative disorders e.g. leukaemia, colorectal, lung, breast
CC and other cancers. The nucleic acid coding for CACNA1G is useful as a
CC marker for screening cancer and age related diseases. A diagnostic kit
CC containing primers (AAS01574-AAS01623) for amplification of a
CC CpG-containing nucleic acid, where the primer hybridises with a target
CC polynucleotide sequence (AAS01627-AAS01676), can be used for detecting
CC aberrant methylation. The CpG island sequences (AAS01677-AAS01692) are
CC selected from genes encoding CACNA1G, apolipoprotein B (APOB), caudal
CC type homeobox transcription factor 2 (CDX2), epidermal growth factor
CC receptor (EGFR), fibrillin-1 (FBN1), G protein-coupled receptor 37
CC (GPR37), heat shock 70kD protein 6 (HSP70B; HSPA6), IQGAP2, Klotho (KL),
CC proteinase-activated receptor 2 (PAR2), paired-like homeodomain
CC transcription factor 2 (PTRX2), patched A and B (PTCHA; PTCHB) and
CC syndecan 1 and 4 (SDC1; SDC4) or a MINT31 sequence.
XX
XX Sequence 22 BP; 15 A; 3 C; 0 G; 4 T; 0 other;
SQ
Query Match 4.38; Score 14; DB 22; Length 22;
Best Local Similarity 100.0%; Pred. No. 6,8e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 196 ttattttatttta 209
Db 21 ttattttatttttta 8

```

```

RESULT 19
AAS01643/c
ID AAS01643 standard; DNA: 22 BP.
XX
XX AAS01643;
AC
XX
XX 18-JUL-2001 (first entry)
DT
XX
XX Human IQGAP2 5'-target sequence for bisulfite PCR.
DE
XX
XX Human: T-type calcium channel; CACNA1G; cytosine methylation; CpG island;
XX cellular proliferative disorder; colorectal cancer; age related disease;
XX apolipoprotein B; APOB; caudal type homeobox transcription factor 2;
XX CDX2; epidermal growth factor receptor; EGFR; fibrillin-1; FBN1;
XX G protein-coupled receptor 37; GPR37; heat shock 70kD protein 6; HSP70B;
XX HSPA6; RasGAP-related protein; IQGAP2; proteinase-activated receptor 2;
XX PAR2; paired-like homeodomain transcription factor 2; PTRX2; Klotho; KL;
XX patched A; patched B; PTCHA; PTCHB; syndecan 1; syndecan 4; SDC1; SDC4;
XX chromosome 5q; ds.
KM

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RESULT 20
AAV07952/c
ID AAV07952 standard; DNA: 26 BP.
XX
XX AAV07952;
AC
XX
XX 02-FEB-1999 (first entry)
DT
XX
XX Helicobacter pylori polypeptide GHPD 1414 5' DNA primer.
DE
XX
XX GHPD 1414; infection; gastritis; ulcer; vaccine; diagnosis;
XX therapy; PCR; primer; ss.
XX
XX Synthetic.
OS Helicobacter pylori.

```

XX MO9843479-A1.
PN 08-OCT-1998.
PD
XX 31-MAR-1998; 98WO-US06421.
PF
XX 01-APR-1997; 97US-0834666.
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PI (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX WPI; 1998-568251/48.
DR
XX
XX
PT New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
XX Claim 5; Page 145; 184pp; English.
PS
XX This 5' primer is used with a 3' primer (see AAV07954) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV07921) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73032) designated GHPO 1414.
CC The isolated polynucleotide, and encoded polypeptide, can be used
CC to develop vaccines for the treatment and prevention of Helicobacter
CC infections.
CC
SQ Sequence 26 BP; 14 A; 5 C; 4 G; 3 T; 0 other;

Query Match 4.3%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.7e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 307 ttltcatgtttct 320
Db 18 TTTTCATGTTTCT 5

RESULT 21
AAV07922/C
ID AAV07922 standard; DNA; 26 BP.
XX
XX AAV07922;
AC
XX 02-FEB-1999 (first entry)
DT
XX Helicobacter pylori polypeptide GHPO 386 5' DNA primer.
DE
XX GHPO 386; infection; gastritis; ulcer; vaccine; diagnosis; therapy;
KW PCR; primer; ss.
XX
XX Synthetic.
OS Helicobacter pylori.
XX
XX MO9843479-A1.
PN
XX 08-OCT-1998.
PD
XX 31-MAR-1998; 98WO-US06421.
PF
XX 01-APR-1997; 97US-0834666.
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PI (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX

DR WPI; 1998-568251/48.
XX
XX
PT New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
XX Claim 5; Page 137; 184pp; English.
PS
XX This 5' primer is used with a 3' primer (see AAV07924) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV72001
CC) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73022) designated GHPO 386.
CC The isolated polynucleotide, and encoded polypeptide, can be used to
CC develop vaccines for the treatment and prevention of Helicobacter
CC infections.
CC
SQ Sequence 26 BP; 15 A; 5 C; 4 G; 2 T; 0 other;

Query Match 4.3%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.7e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 307 ttltcatgtttct 320
Db 20 TTTTCATGTTTCT 7

RESULT 22
AAV07937/C
ID AAV07937 standard; DNA; 27 BP.
XX
XX AAV07937;
AC
XX 02-FEB-1999 (first entry)
DT
XX Helicobacter pylori polypeptide GHPO 896 5' DNA primer.
DE
XX GHPO 896; infection; gastritis; ulcer; vaccine; diagnosis;
KW Therapy; PCR; primer; ss.
XX
XX Synthetic.
OS Helicobacter pylori.
XX
XX MO9843479-A1.
PN
XX 08-OCT-1998.
PD
XX 31-MAR-1998; 98WO-US06421.
PF
XX 01-APR-1997; 97US-0834666.
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PI (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX WPI; 1998-568251/48.
DR
XX
XX
PT New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
XX Claim 5; Page 141; 184pp; English.
PS
XX This 5' primer is used with a 3' primer (see AAV07939) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV07916) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73027) designated GHPO 896.
CC The isolated polynucleotide, and encoded polypeptide, can be used
CC to develop vaccines for the treatment and prevention of Helicobacter

CC infections.
 XX
 SQ Sequence 27 BP; 14 A; 5 C; 4 G; 4 T; 0 other;

Query Match 4.3%; Score 14; DB 19; Length 27;
 Best Local Similarity 100.0%; Pred. No. 6.7e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 307 ttttcagttttct 320
 |||||||
 Db 18 ttttcagttttct 5

RESULT 23
 AAT42655/c
 ID AAT42655 standard; DNA; 29 BP.
 XX
 AC AAT42655;
 XX
 DT 25-FEB-1997 (first entry)
 XX
 DE Primer for amplifying verotoxin (VT-1) subunit A coding sequence.
 XX
 KM Verotoxin; Escherichia coli; enteric infection; diarrhoea; vaccine;
 KM haemolytic uraemic syndrome; detection; ss.
 XX
 OS Synthetic.
 XX
 PN M09630043-A1.
 XX
 PD 03-OCT-1996.
 XX
 PF 25-MAR-1996; 96MO-US04093.
 XX
 PR 24-MAR-1995; 95US-0410058.
 XX
 PA (OPHI-) OPHIDIAN PHARM INC.
 XX
 PI Carroll SB, Padhye NV, Stafford DC;
 XX
 DR WPI; 1996-505779/50.
 XX
 PT Compn. contg. neutralising antitoxin against E.coli vero-toxin -
 PT used to treat intoxicated individuals, and as a prophylactic against
 PT diarrhoeal disease or extra-intestinal complications of E.coli
 PT infection
 XX
 PS Example 6; Page 58; 101pp; English.
 XX
 CC Compositions containing neutralising antitoxin against one or more E.
 CC coli verotoxin (VT) can be used to treat intoxicated adults and
 CC children with enteric bacterial infections. They may also be used as
 CC prophylactics e.g. as a vaccine, against diarrhoeal disease or the
 CC development of extra-intestinal complications of E.coli infection.
 CC especially haemolytic uraemic syndrome. The antitoxin can also be
 CC used to detect E. coli VT in a sample. The VT is recombinant,
 CC preferably a fusion protein containing a non-VT protein sequence and
 CC part of the E.coli VT1 or VT2 sequence. Two primers (AAT42655,
 CC AAT42656) were used to amplify the verotoxin VT-1 A subunit coding
 CC sequence and add a histidine tag coding sequence to the subunit
 CC sequence. Two primers (AAT42655, AAT42658) were used to amplify the
 CC verotoxin VT-1 A and B subunits and add a histidine tag coding
 CC sequence to the subunit sequences.
 XX
 SQ Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.3%; Score 14; DB 17; Length 29;
 Best Local Similarity 100.0%; Pred. No. 6.6e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 189 aaataattattt 202

Db 22 |||||||
 22 AAATATTATT 9

RESULT 24
 AAA51200/c
 ID AAA51200 standard; DNA; 29 BP.
 XX
 AC AAA51200;
 XX
 DT 26-SEP-2000 (first entry)
 XX
 DE N-terminal primer for E. coli verotoxin 1 subunit A gene.
 XX
 KM VT-1; verotoxin; antitoxin therapy; fusion protein; affinity tag; food;
 KM recombinant production; screening; dairy; anti-bacterial; vaccine;
 KM primer; polystyridine; ss.
 XX
 OS Escherichia coli.
 OS Synthetic.
 OS
 PN US6080400-A.
 XX
 PD 27-JUN-2000.
 XX
 PF 13-MAR-1997; 97US-0816977.
 XX
 PR 24-MAR-1995; 95US-0410058.
 XX
 PA (OPHI-) OPHIDIAN PHARM INC.
 XX
 PI Williams JA, Byrne LM;
 XX
 DR WPI; 2000-451195/39.
 XX
 PT Bacterial cell for recombinantly expressing bacterial toxins in large
 PT quantities useful for immunization and treatment of bacterial
 PT infections, comprises expression vector encoding bacterial toxin
 XX
 PS Example 6; Column 83; 83pp; English.
 XX
 CC E. coli verotoxin (VT) type 1 and 2 subunits A and B were cloned into
 CC pET-23b, designed to allow expression of the native proteins containing
 CC C-terminal polystyridine tags. The VT-1 and VT-2 genes were engineered
 CC to convert the signal sequence methionine codon into a NdeI site to
 CC allow cloning of the amplified genes into the vector without addition of
 CC vector-encoded amino acids. The C-terminal primers comprises the
 CC C-terminal 7 codons of each gene fused to the sequence CTCGAGCC, in order
 CC to add the polystyridine tag. The primers delete the native stop codons,
 CC and when cloned into pET-23 add a C-terminal extension of Leu-Glu-(His)6.
 CC VT B chains are small proteins (approximately 8 kDa), so use of a small
 CC affinity tag was preferred (i.e. polystyridine). A polystyridine affinity
 CC tag facilitates single step affinity purification of subunits from
 CC periplasmic extracts. However, due to poor recovery of his-tagged VT-1 A
 CC and VT-2 A chains, expression of maltose binding protein (MBP) fused
 CC subunits was undertaken. Due to the toxicity of the VT-2 B subunit,
 CC strict uninduced promoter control is necessary to permit cell viability.
 CC Bacterial host cells expressing a recombinant expression vector encoding
 CC a polystyridine affinity tag and a portion of the VT-2 B chain are
 CC claimed. The vector is chosen from pET24hisVT2BL+, pET24hisVT2BL- and
 CC pET24VT2B, where "L+" indicates that the vector encodes the preprotein
 CC form of the protein and "L-" indicates that the vector encodes the mature
 CC form of the protein. The bacterial cell is capable of expressing large
 CC quantities (40 mg/l) of VT-2B. The toxins are useful for immunizing
 CC non-mammals and for detecting bacterial toxins in environmental samples
 CC including soil, water, industrial samples, biological samples and samples
 CC obtained from food and dairy processing instruments.
 XX
 SQ Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 29;
 Best Local Similarity 100.0%; Pred. No. 6.6e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 189 aaataattatttt 202
 |||||
 Db 22 AAAATAATTATTTT 9

RESULT 25

AAAT25703/C
 ID AAT25703 standard; cDNA to mRNA; 31 BP.

AC AAT25703;

DT 10-OCT-1996 (first entry)

DE Human gene signature HUNGSO7904.

KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;
 human; cloning; mapping; non-biased library; diagnosis; detection;
 cell typing; abnormal cell function; ss.

OS Homo sapiens.

PN WO9514772-A1.

PD 01-JUN-1995.

PF 11-NOV-1994; 94WO-JP01916.

PR 12-NOV-1993; 93JP-0355504.

PA (MATS/) MATSUBARA K.

PA (OKUB/) OKUBO K.

PI Matsubara K, Okubo K;

DR WPI; 1995-206931/27.

PT Identifying gene signatures in 3'-directed human cDNA library - e.g.
 for diagnosis of abnormal cell function, by preparing cDNA that
 reflects relative abundance of corresp. mRNA in specific human
 tissues

PS Claim 1; Page 1910; 2245pp; Japanese.

CC A single-stranded DNA (or its complementary strand or the corresp.
 double-stranded DNA) which comprises one of the 7837 "GS" sequences
 given in AAT19001-T26837 and which is able to hybridise to part of
 human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
 sequences were obtained from 3'-directed cDNA libraries prepared
 from various human tissues: synthesis of cDNA was initiated from the
 3'-end of mRNA by using poly(1) as the sole primer. Since the 3'-
 untranslated sequence is unique to a particular mRNA species, almost
 all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
 is constructed so as to reflect accurately the relative abundance of
 different mRNAs in the particular tissue from which it was derived.
 CC The appearance frequency of a given GS in a cDNA library can be
 determined (esp. using primers and probes derived from the GS
 sequences) as a means of diagnosing abnormal cell function or for
 recognising different cell types.

CC Sequence 31 BP; 14 A; 3 C; 3 G; 10 T; 1 other;

Query Match 4.3%; Score 14; DB 16; Length 31;
 Best Local Similarity 100.0%; Pred. No. 6.6e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 198 attttattttaa 211
 |||||

Db 20 ATTGTTATTTTAA 7

RESULT 26
 AAV67854
 ID AAV67854 standard; DNA; 31 BP.

AC AAV67854;

DT 24-DEC-1998 (first entry)

DE Nucleotide fragment containing polymorphic site, WI-11163.

KW ss; polymorphic site; nucleic acid analysis; diagnosis; monitoring;
 cancer; inflammation; heart disease; CNS disease.

OS Homo sapiens.

PN WO9838846-A2.

PD 11-SEP-1998.

PF 06-MAR-1998; 98WO-US04571.

PR 28-MAR-1997; 97US-0042125.

PR 07-MAR-1997; 97US-0813159.

PA (AFFY-) AFFYMETRIX INC.

PI Berno A, Chee M, Fan J, Lipschutz RJ;

DR WPI; 1998-495419/42.

PT New nucleic acid segments containing polymorphic sites, or
 complements and methods of detecting a nucleic acid - for general
 use including diagnosis and monitoring of diseases

PS Claim 1; Page 25; 42pp; English.

CC New nucleic acid segment comprising one of the 10 - 100 bp sequences
 given in the specification (sequences of a polymorphic site), or the
 complement of the segment and a method of analysing a nucleic acid
 comprising determining the base occupying the polymorphic site of the
 CC polymorphic fragment sequences are disclosed in the specification. The
 CC information obtained from nucleic acid analysis by the method described
 CC is useful in diagnosis or monitoring of diseases like cancer,
 CC inflammation, heart disease, CNS diseases, and susceptibility to
 CC infection by microorganisms. In addition, the nucleic acid segments are
 CC useful in manufacturing medication in the treatment of prophylaxis of
 CC diseases, and also the use of the DNA segments as pharmaceutical.

CC Sequence 31 BP; 17 A; 2 C; 4 G; 7 T; 1 other;

Query Match 4.3%; Score 14; DB 19; Length 31;
 Best Local Similarity 100.0%; Pred. No. 6.6e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 209 aaattgtttaaaa 222
 |||||

Db 17 aaattgtttaaaa 30

RESULT 27

AAZ27689/C
 ID AAZ27689 standard; DNA; 32 BP.

AC AAZ27689;

DT 22-DEC-1999 (first entry)

DE PCR primer for Verotoxin gene.

KW Verotoxin; VT1; VT2; detection; PCR primer; ss.

OS Synthetic.

```
OS Escherichia coli.
XX JP11243996-A.
XX
XX PD 14-SEP-1999.
XX PF 27-FEB-1998; 98UP-0047677.
XX PR 27-FEB-1998; 98UP-0047677.
XX PA (TOYOBO ) TOYOBOKK.
XX DR WPI; 1999-603716/52.
XX PT An oligonucleotide for amplification of verotoxin - useful in the
PT detection of inactivated verotoxin gene by transfer of a foreign DNA
PS fragment
PS Claim 11; Page 9; 10pp; Japanese.
XX
XX This sequence represents a PCR primer of the invention. The primer is
CC used for amplification of the E. coli verotoxin (VT) gene. The
CC oligonucleotide is useful for detection of inactivated VT gene by
CC transfer of a foreign DNA fragment. Simple, rapid and specific
CC amplification of VT gene from environmental factors is achieved using the
CC oligonucleotide of the invention.
XX
XX Sequence 32 BP; 12 A; 2 C; 4 G; 14 T; 0 other;
SO
OY Query Match 4.3%; Score 14; DB 20; Length 32;
Best Local Similarity 100.0%; Pred. No. 6.6e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0
DB 189 aaataattatctt 202
17 AAATAATTATTTT 4
RESULT 28
AAAT97603/C
ID AT97603 standard; DNA; 36 BP.
XX AC AT97603;
XX DT 30-APR-1998 (first entry)
XX DE Shigella dysenteriae delta-stx allele PCR primer 13.
XX KW Delta-virg allele; delta-guaB-A allele; PCR; amplification; primer;
XX delta-stx allele; shigellosis; vaccine; ss.
XX OS Synthetic.
XX OS Shigella dysenteriae.
XX PN WO9737685-A1.
XX PD 16-OCT-1997.
XX PF 09-APR-1997; 97WO-US05954.
XX PR 09-APR-1996; 96US-0629600.
XX PA (UYMA-) UNIV MARYLAND BALTIMORE.
XX PI Levine MM, Noriega FR;
XX WPI; 1997-512417/47.
XX DR Shigella mutants with mutation in guaB-A - used in vaccines against
XX Shigellosis
PS Example 6; Page 57; 94pp; English.
```

Query Match	Best Local Similarity	4.3%;	Score 14;	DB 18;	Length 36;
Matches	14;	Conservative	0;	Mismatches	0;
				Indels	0;
				Gaps	0;
189 aaataattattt 202	27 AAAATATATTATT 14				
<p> This is a PCR primer used in the amplification of the Shigella dysenteriae 1 delta-stx allele. The delta-stx allele was integrated into delta-guab-A of delta-guab-A, delta-virg S. dysenteriae 1, which inactivated the shiga toxin of this strain. The mutant can be used in the preparation of vaccines such as a live vector vaccine comprising a Shigella mutant, (which encodes and expresses a foreign antigen, and a pharmaceutically acceptable carrier) or a DNA mediated vaccine comprising the Shigella mutant (which also contains a plasmid which encodes and expresses a foreign antigen in a eukaryotic cell). The vaccines can be used against Shigellosis. </p>					
Sequence 36 BP; 11 A; 3 C; 10 G; 12 T; 0 other;					
<p> RESULT 29 AAC90606 AAC90606 standard; RNA; 36 BP. </p>					
<p> AAC90606; 20-MAR-2001 (first entry) </p>					
<p> Tomato spotted wilt virus S RNA partial sequence #10. </p>					
<p> Tospovirus resistance; transgenic plant; tomato spotted wilt virus; Impatiens necrotic spot virus; TSMV; ss. </p>					
<p> Tomato spotted wilt virus. </p>					
<p> Key Location/Qualifiers misc_binding 1..13 /tag= a /bound_moiety= "binds nucleotides 33-21 of AAC89655" misc_binding 15..28 /tag= b /bound_moiety= "binds nucleotides 19-6 of AAC89655" misc_binding 32..36 /tag= C /bound_moiety= "binds nucleotides 5-1 of AAC89655" </p>					
<p> US6150585-A. 21-NOV-2000. </p>					
<p> 26-NOV-1996; 96US-0757011. </p>					
<p> 02-MAY-1991; 91US-0694734. 14-APR-1993; 93US-0047346. 26-OCT-1993; 93US-0143397. 27-JUL-1994; 94US-0280903. 03-NOV-1994; 89US-0431259. 05-DEC-1989; 89US-0446024. </p>					
<p> (NOVS.) NOVARTIS FINANCE CORP. Peters D, Gielen J, De Haan P, Van Grinsven M, Kool A; Goldbach RW; WPI: 2001-060031/07. </p>					
<p> Recombinant DNA construct comprising a DNA sequence encoding an RNA sequence that codes for a tospovirus protein, useful for producing plants with reduced susceptibility to tospovirus infection - </p>					

XX Example 9; Fig 16C; 49pp; English.

CC The present invention provides DNA constructs encoding RNA sequences from
CC a tospovirus which can be used to produce transgenic plants with immunity
CC to tospoviruses. Examples of tospoviruses include the tomato spotted wilt
CC virus and the Impatiens necrotic spot virus.

SO Sequence 36 BP; 7 A; 0 C; 0 G; 29 U; 0 other;

Query Match 4.3%; Score 14; DB 22; Length 36;

Best Local Similarity 21.4%; Pred. No. 6.5e+03;

Matches 3; Conservative 11; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttta 209

|||||:|||||

DB 7 uuuuuuuuuuuua 20

RESULT 30

AA062952

ID AA062952 standard; DNA; 37 BP.

XX AA062952;

DT 09-SEP-1994 (first entry)

XX Glycophorin antibody 1C3 Fab coding region PCR primer.

XX Glycophorin; antibody 1C3; target binding polypeptide; PCR;

KW polymerase chain reaction; primer: antibody engineering;

KW humanized antibody; phagemid pHEA; plasmid p56;ss.

XX Synthetic.

XX WO9407921-A.

PD 14-APR-1994.

XX 24-SEP-1993; 93WO-AU00491.

XX 25-SEP-1992; 92AU-0004973.

PA (CSIR) COMMONWEALTH SCI & IND RES ORG.

PI Atwell JL, Colman PM, Hudson PJ, Irving RA, Kortt A;

PI Lah M, Malbyrl, Power BE;

DR WPI; 1994-135515/16.

XX New target-binding polypeptide(s) used for diagnosis, etc.

PT Having a stable core polypeptide region with at least one

PT target-binding region covalently attached, opt. mutated to alter

PT specificity, etc.

PS Example; Page 36; 67pp; English.

XX PCR primers given in AA062951-52 were used to clone anti-glycophorin

CC antibody 1C3 Fab coding region. The DNA sequence of the first 1443

CC bases of the Fab fragment in pHEA, ready for ligation post PCR

CC amplification for ligation into p569, is given in AA062958.

XX Sequence 37 BP; 14 A; 5 C; 0 G; 18 T; 0 other;

SO Query Match 4.3%; Score 14; DB 15; Length 37;

Best Local Similarity 100.0%; Pred. No. 6.5e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 31

AAF55449/C

ID AAF55449 standard; DNA; 45 BP.

XX AAF55449;

DT 29-MAY-2001 (first entry)

XX Oligonucleotide used to construct the shuttle vector pAAO-E-TATA.

XX Adenovirus vector; gene delivery vector; E2B gene; E4 gene; vaccine;

KW replication defective virus; cell proliferation; cell differentiation;

KW gene therapy; ss.

XX Synthetic.

XX EPI083229-A1.

PD 14-MAR-2001.

XX 10-SEP-1999; 99EP-0202966.

XX 10-SEP-1999; 99EP-0202966.

PA (INTR-) INTROGENE BV.

XX WPI; 2001-228258/24.

XX Producing a recombinant adenovirus-like gene delivery vehicle with

PT modified E2B or E4 functions, for gene therapy, comprises generating an

PT adenoviral vector where E2B or E4 is under the control of a synthetic

PT promoter

XX Example 1; Page 10; 56pp; English.

XX The specification describes a method for producing a recombinant

CC adenovirus-like gene delivery vehicle having reduced expression of

CC adenoviral E2B and/or E4 gene products in a target cell. The method

CC comprises generating a recombinant adenoviral vector lacking E1A and

CC E1B sequences, but having at least the E2B and/or E4 sequences encoding

CC products essential for adenoviral replication. Compared with previous

CC vectors, the new recombinant vectors are replication defective and

CC express the remaining viral genes only at background levels. The vector

CC itself does not dominantly elicit a response of the immune system, but

CC the immune response is directed primarily against the transgene product,

CC and is less toxic to the cells which, in turn, results in a prolonged

CC synthesis of the protein of interest. The recombinant adenoviral

CC vectors are used in the functional characterization of gene products in

CC cells, tissues or animals in order to find genes that encode for

CC proteins with a desired function such as those that interfere with cell

CC proliferation and differentiation. The vectors are further used as

CC vaccines, in gene therapy, and for protein production in mammalian

CC cells. Oligonucleotides AAF55449-50 were annealed together, and used to

XX construct a shuttle vector which was used in the method of the invention.

SO Sequence 45 BP; 14 A; 5 C; 7 G; 19 T; 0 other;

Query Match 4.3%; Score 14; DB 22; Length 45;

Best Local Similarity 100.0%; Pred. No. 6.4e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92

|||||:|||||

DB 21 CTGAAATTATTA 8

RESULT 32

AAF55450

ID AAF55450 standard; DNA; 45 BP.

XX	AA655450;	
XX		
DT	29-MAY-2001	(first entry)
XX		
DE	Oligonucleotide used to construct the shuttle vector PMAO-E-TATA.	
XX		
KM	Adenovirus vector; gene delivery vector; E2B gene; E4 gene; vaccine;	
KW	replication defective virus; cell proliferation; cell differentiation;	
XX	gene therapy; ss.	
XX		
OS	Synthetic.	
XX		
PM	EP1083229-A1.	
XX		
PD	14-MAR-2001.	
XX		
PF	10-SEP-1999;	99EP-0202966.
XX		
PR	10-SEP-1999;	99EP-0202966.
XX		
PA	(INTR-) INTROGENE BV.	
XX		
DR	WPI, 2001-228258/24.	
XX		
PT	Producing a recombinant adenovirus-like gene delivery vehicle with	
PT	modified E2B or E4 functions, for gene therapy, comprises generating an	
PT	adenoviral vector where E2B or E4 is under the control of a synthetic	
PT	promoter	
XX		
PS	Example 1; Page 10; 56pp; English.	
XX		
CC	The specification describes a method for producing a recombinant	
CC	adenovirus-like gene delivery vehicle having reduced expression of	
CC	adenoviral E2B and/or E4 gene products in a target cell. The method	
CC	comprises generating a recombinant adenoviral vector lacking E1A and	
CC	E1B sequences, but having at least the E2B and/or E4 sequences encoding	
CC	products essential for adenoviral replication. Compared with previous	
CC	vectors, the new recombinant vectors are replication defective and	
CC	express the remaining viral genes only at background levels. The vector	
CC	itself does not dominantly elicit a response of the immune system, but	
CC	the immune response is directed primarily against the transgene product,	
CC	and is less toxic to the cells which, in turn, results in a prolonged	
CC	synthesis of the protein of interest. The recombinant adenoviral	
CC	vectors are used in the functional characterization of gene products	
CC	in vitro or in vivo, and for overexpression of known and novel genes in	
CC	cells lines, tissues or animals in order to find genes that encode for	
CC	proteins with a desired function such as those that interfere with cell	
CC	proliferation and differentiation. The vectors are further used as	
CC	vaccines, in gene therapy, and for protein production in mammalian	
CC	cells. Oligonucleotides AA655449-50 were annealed together, and used to	
CC	construct a shuttle vector which was used in the method of the invention.	
XX		
50	Sequence 45 BP; 19 A; 7 C; 5 G; 14 T; 0 other;	
XX		
DT	Query Match	4.3%; Score 14; DB 22; Length 45;
DT	Best Local Similarity	100.0%; Pred. No. 6.4e+03;
DT	Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
OY	79 ctgaataattatcaa 92	
DB	29 ctgaataattatcaa 42	
XX		
RESULT 33		
XX	AA655449/c	
XX	AA655449 standard; DNA; 45 BP.	
XX	AA655449;	
XX		
DT	05-MAR-2001	(first entry)
DT		

Query Match	Best Local Similarity	4.3%;	Score 14;	DB 22;	Length 45;
Matches	14;	Conservative	0;	Mismatches	0;
Indels	0;	Gaps	0;		
Qy	79	ctgaataattatca	92		
Db	21	CTGAATAATTATTA	8		
RESULT	34				
AAC88875					
ID	AAC88875	standard;	DNA;	45	BP.
AC	AAC88875;				
DT	05-MAR-2001	(first entry)			
DE	Oligonucleotide	TATAmn.			
KW	Adenovirus type 35;	Ad35;	adenovirus type 5;	Ad5;	gene delivery vehicle;
OS	gene therapy;	linker;	ss.		
PN	Synthetic.				
PI	EPI054064-A1.				
PD	22-NOV-2000.				
PF	16-MAY-2000;	2000EP-0201738.			
PR	17-MAY-1999;	99EP-0201545.			
PA	(INTR-)	INTROGENE BV.			
PI	Bout A,	Vogels R,	Haveinga MJE;		
DR	WPI;	2001-001097/01.			
CC	Adenovirus derived gene delivery vehicles comprising at least one				
CC	element of adenovirus type 35, efficiently transfers genetic material				
CC	to a human cell without the inherent problem of pre-existing immunity -				
CC	Example 13; Page 31; 138pp; English.				
CC	The present sequence is a linker used in a gene delivery vehicle				
CC	comprising an element of adenovirus type 35 or a functional equivalent				
CC	of such an element. The element is responsible for avoiding or reducing				
CC	neutralising activity against adenoviral elements by the host to which				
CC	the gene is to be delivered. The vehicle can be used to deliver genes or				
CC	nucleic acids of interest to host cells. Use of the delivery system				
CC	efficiently transfers genetic material to a human cell without the				
CC	inherent problem of pre-existing immunity, found with previous serotypes.				
CC	Sequence 45 BP; 14 A; 5 C; 7 G; 19 T; 0 other;				

PI Bout A, Vogels R, Havenga MTE;
XX
DR WPI: 2001-001097/01.
XX
PR Adenovirus derived gene delivery vehicles comprising at least one
XX element of adenovirus type 35, efficiently transfers genetic material
XX to a human cell without the inherent problem of pre-existing immunity -
XX
PS Example 13; Page 31; 138pp; English.
XX
CC The present sequence is a linker used in a gene delivery vehicle
CC comprising an element of adenovirus type 35 or a functional equivalent
CC of such an element. The element is responsible for avoiding or reducing
CC neutralising activity against adenoviral elements by the host to which
CC the gene is to be delivered. The vehicle can be used to deliver genes or
CC nucleic acids of interest to host cells. Use of the delivery system
CC efficiently transfers genetic material to a human cell without the
CC inherent problem of pre-existing immunity, found with previous serotypes.
CC
SQ Sequence 45 BP; 19 A; 7 C; 5 G; 14 T; 0 other;

QY 79 ctgaattattataa 92
DB 29 ctgaattattataa 42

RESULT 35
AAZ66366
ID AAZ66366 standard; DNA; 47 BP.
XX
AC AAZ66366;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human map-related biallelic marker SEQ ID NO:713.
XX
KW Human genome; biallelic marker; high density disequilibrium map;
KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW haplotyping; hybridisation; identification; characterisation;
KW diagnosis; single nucleotide polymorphism; SNP; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(24,C)
FT /*tag= a
FT /standard_name="single nucleotide polymorphism"
XX
PN WO9954500-A2.
XX
PD 28-OCT-1999.
XX
PE 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
PR 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
XX
DR WPI: 2000-013267/01.
XX
PT Novel biallelic markers used to construct a high density disequilibrium
XX map of the human genome -
XX
PS Claim 1; Page 382; 2745pp; English.
XX

CC AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their
CC nucleotide sequences. AAZ6579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the
CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and
CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the sequence listing
CC from the present invention.
XX
SQ Sequence 47 BP; 22 A; 2 C; 6 G; 17 T; 0 other;

QY 109 gtaataataataa 122
DB 5 gtaataataataa 18

RESULT 36
AAZ67473/C
ID AAZ67473 standard; DNA; 47 BP.
XX
AC AAZ67473;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human map-related biallelic marker SEQ ID NO:1820.
XX
KW Human genome; biallelic marker; high density disequilibrium map;
KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW haplotyping; hybridisation; identification; characterisation;
KW diagnosis; single nucleotide polymorphism; SNP; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(24,C)
FT /*tag= a
FT /standard_name="single nucleotide polymorphism"
XX
PN WO9954500-A2.
XX
PD 28-OCT-1999.
XX
PE 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
PR 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
XX
DR WPI: 2000-013267/01.
XX
PT Novel biallelic markers used to construct a high density disequilibrium
XX map of the human genome -
XX
PS Claim 1; Page 618; 2745pp; English.
XX
CC AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their

CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the
CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and
CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.

XX
SQ Sequence 47 BP; 16 A; 4 C; 2 G; 25 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 6.4e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 162 atataataaatt 175
|||||
DB 23 ATATATAATAAATT 10

RESULT 37
AAZ67533/C
ID AAZ67533 standard; DNA: 47 BP.
XX
AC AAZ67533;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human map-related biallelic marker SEQ ID NO:1880.
XX
KW Human genome; biallelic marker; high density disequilibrium map;
KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW haplotyping; hybridisation; identification; characterisation;
KW diagnosis; single nucleotide polymorphism; SNP; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(24,C)
FT /*tag= a
FT /standard_name= "single nucleotide polymorphism"
XX
PN WO9954500-A2.
XX
PD 28-OCT-1999.
XX
PF 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
PR 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
XX
DR WPI: 2000-013267/01.
XX
PT Novel biallelic markers used to construct a high density disequilibrium
PT map of the human genome -
XX
PS Claim 1; Page 631; 2745pp; English.
XX
CC AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their
CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the

CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and
CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.

XX
SQ Sequence 47 BP; 15 A; 6 C; 6 G; 20 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 6.4e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 95 atatgataagtata 108
|||||
DB 30 ATATGATAAGTATA 17

RESULT 38
AAZ67549/C
ID AAZ67549 standard; DNA: 47 BP.
XX
AC AAZ67549;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human map-related biallelic marker SEQ ID NO:1896.
XX
KW Human genome; biallelic marker; high density disequilibrium map;
KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW haplotyping; hybridisation; identification; characterisation;
KW diagnosis; single nucleotide polymorphism; SNP; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(24,G)
FT /*tag= a
FT /standard_name= "single nucleotide polymorphism"
XX
PN WO9954500-A2.
XX
PD 28-OCT-1999.
XX
PF 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
PR 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
XX
DR WPI: 2000-013267/01.
XX
PT Novel biallelic markers used to construct a high density disequilibrium
PT map of the human genome -
XX
PS Claim 1; Page 635; 2745pp; English.
XX
CC AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their
CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the
CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and

CC haplotyping studies which are useful in determining the genetic basis
 CC for disease states. Compositions and methods of the invention can also
 CC be useful for the identification of the targets for the development of
 CC pharmaceutical agents and diagnostic methods, as well as the
 CC characterisation of the differential efficacious responses to and side
 CC effects from pharmaceutical agents acting on a disease as well as other
 CC treatment.
 CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
 CC and 3367, are not actually given a sequence in the Sequence Listing
 CC from the present invention.
 CC
 SQ Sequence 47 BP; 17 A; 10 C; 2 G; 18 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 47;
 Best Local Similarity 100.0%; Pred. No. 6.4e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 148 ttttaaaaggaaa 161
 ||||||||||||
 Db 27 TTTTAAAAAGCAA 14

RESULT 39

AAZ67813
 ID AAZ67813 standard; DNA; 47 BP.
 XX
 AC AAZ67813;
 XX
 DT 10-SEP-2001 (first entry)
 XX

DE Human map-related diallelic marker SEQ ID NO:2160.
 XX
 KW Human genome; diallelic marker; high density disequilibrium map;
 KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
 KW haplotyping; hybridisation; identification; characterisation;
 KW diagnosis; single nucleotide polymorphism; SNP; ds.
 XX
 OS Homo sapiens.
 XX

Key Location/Qualifiers
 FT variation replace(24,G)
 FT /*tag=a
 FT /standard_name="single nucleotide polymorphism"

W0954500-A2.

28-OCT-1999.

21-APR-1999; 99WO-IB00822.

21-APR-1998; 98US-0082614.
 PR 23-NOV-1998; 98US-0109732.

(GEST) GENSET.

Cohen D, Blumenfeld M, Chumakov I;

WPI: 2000-013267/01.

Novel diallelic markers used to construct a high density disequilibrium
 map of the human genome -

Claim 1; Page 691; 2745pp; English.

AAZ65654 to AAZ69578 represent human diallelic markers from the present
 CC invention, which contain a polymorphic base at position 24 of their
 CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
 CC primers for the diallelic markers. The diallelic markers of the
 CC invention have a variety of uses; they can be used for high density
 CC mapping of the human genome, and in complex association studies and
 CC haplotyping studies which are useful in determining the genetic basis
 CC for disease states. Compositions and methods of the invention can also

CC be useful for the identification of the targets for the development of
 CC pharmaceutical agents and diagnostic methods, as well as the
 CC characterisation of the differential efficacious responses to and side
 CC effects from pharmaceutical agents acting on a disease as well as other
 CC treatment.
 CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
 CC and 3367, are not actually given a sequence in the Sequence Listing
 CC from the present invention.
 CC
 SQ Sequence 47 BP; 19 A; 4 C; 8 G; 16 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 47;
 Best Local Similarity 100.0%; Pred. No. 6.4e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 198 attttatttttaa 211
 ||||||||||||
 Db 11 attttatttttaa 24

RESULT 40

AAA98312
 ID AAA98312 standard; DNA; 50 BP.
 XX
 AC AAA98312;
 XX
 DT 02-FEB-2001 (first entry)
 XX

DE Human MSH6 fragment 8/exon 8 to 10 DNA Ref-Seq fragment.

KW Human mismatch repair gene; hMSH6; disease predisposition; genotype;
 KW mutation; carcinoma; colorectal; endometrial; ovarian; leukemia;
 KW neoplastic disease; drug development; ds.
 XX
 OS Homo sapiens.
 XX

DE19909878-A1.

07-SEP-2000.

06-MAR-1999; 99DE-1009878.

06-MAR-1999; 99DE-1009878.

(UYDR) UNIV DRESDEN TECH.

Plaschke J, Kruppa C, Schackert H;

WPI: 2000-588378/56.

Novel variants of the human mismatch repair gene, MSH6, useful e.g. for
 determining predisposition to cancer and for development of drugs -

Disclosure; Fig 3; 14pp; German.

This invention describes a novel method of determining a predisposition
 CC to disease by genotyping a subject's DNA sequence (A) of the human
 CC mismatch repair gene, MSH6 at specified positions and comparing with
 CC reference DNA sequences, optionally taking into account all possible
 CC combinations of variations of the individual mutations, including any
 CC chosen absolute number of variations. (A), and analysis of their
 CC sequences, are useful for the following: (i) determining a predisposition
 CC to disease, especially colorectal, endometrial and ovarian carcinoma and
 CC leukemia; (ii) determining an increased mutation rate (frequency of base
 CC substitutions, insertions and/or deletions) in eukaryotic cells; (iii)
 CC predicting the progression, severity and survival time of patients with
 CC neoplastic disease; (iv) the development of therapeutic and 'life-style'
 CC drugs; (v) predicting individual differences in response to known
 CC chemotherapeutic agents (e.g. cis-platin) or drugs developed from (iv);
 CC (vi) optimizing individual treatments and interventions against
 CC neoplasia; (vii) controlling the mutation rate in eukaryotic cells, in
 CC vitro or in vivo; (viii) constructing genes and vectors, particularly for

CC development of pharmaceuticals: (ix) developing diagnostic kits and other
 CC systems for genotyping; and (x) developing in vivo and in vitro test
 CC systems for expressing individual forms of the MSH6 gene, e.g. for
 CC studying pathophysiology of disease or processes in which MSH6 is
 CC involved, and for drug development and testing.

XX Sequence 50 BP; 8 A; 8 C; 5 G; 29 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 50;

Best Local Similarity 100.0%; Pred. No. 6.3e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 140 cttttgttttaa 153

Db 11 cttttgttttaa 24

RESULT 41

AAFA8097 standard; DNA; 15 BP.

AAFA8097;

30-MAR-2001 (first entry)

IGFBP3 oligonucleotide #1517.

Antisense therapy: antiproliferative; antiinflammatory; antipsoriatic;
 cytosolic; dermatological; cardiant; virucide; ophthalmological; keloid;
 skin disorder; insulin-like Growth Factor 1 receptor; IGF-1; pityriasis;
 IGF binding protein; IGFBP-2; IGFBP3; inflammation; psoriasis; pilaris;
 growth factor mediated cell proliferation; ichthyosis; serborrhea; ruba;
 keratosis; neoplasia; scleroderma; wart; skin cancer; sclerotic disease;
 hyperneovascular condition; hyperplasia; kidney disease;
 neovascular condition of the retina; ss.

Homo sapiens.

MO200078341-A1.

28-DEC-2000.

21-JUN-2000; 2000MO-AU00693.

21-JUN-1999; 99US-0140345.

(MURD-) MURDOCH CHILDRENS RES INST.

Wraight CJ, Werther GA, Edmondson SR;

WPI; 2001-041421/05.

Ameliorating the effects of a disorder, e.g. psoriasis, by
 administering UV (ultra-violet) treatment (optional) and an antisense
 nucleic acid that inhibits or reduces growth factor mediated cell
 proliferation and/or inflammation -

Example 7; Page 54; 201pp; English.

The present invention relates to a method for ameliorating the effects
 of skin disorders. The method comprises contacting the skin with an
 antisense oligonucleotide, (for Insulin-like Growth Factor [IGF]-1
 receptor, IGF binding protein [IGFBP]-2 or IGFBP3), which is capable of
 inhibiting or reducing growth factor mediated cell proliferation,
 inflammation and/or other disorders. The present sequence is an
 oligonucleotide which can be used to design the antisense
 oligonucleotides of the present invention (see AAF45151 and
 AAF45153-F45161). The method is useful for ameliorating the effects of
 psoriasis, ichthyosis, pityriasis, ruba, pilaris, serborrhea, keloids,
 keratosis, neoplasias, scleroderma, warts, benign growths, cancers of the
 skin, a hyperneovascular condition such as a neovascular condition of the
 retina, brain or skin, growth factor-mediated malignancies, other

CC sclerotic disease, kidney disease, hyperproliferation of the inside of
 CC blood vessels or any other hyperplasia.

XX Sequence 15 BP; 7 A; 1 C; 3 G; 4 T; 0 other;

Query Match 4.0%; Score 13; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 26 agaagaatgttta 38

Db 3 agaagaatgttta 15

RESULT 42

AAFA8098 standard; DNA; 15 BP.

AAFA8098;

30-MAR-2001 (first entry)

IGFBP3 oligonucleotide #1518.

Antisense therapy: antiproliferative; antiinflammatory; antipsoriatic;
 cytosolic; dermatological; cardiant; virucide; ophthalmological; keloid;
 skin disorder; insulin-like Growth Factor 1 receptor; IGF-1; pityriasis;
 IGF binding protein; IGFBP-2; IGFBP3; inflammation; psoriasis; pilaris;
 growth factor mediated cell proliferation; ichthyosis; serborrhea; ruba;
 keratosis; neoplasia; scleroderma; wart; skin cancer; sclerotic disease;
 hyperneovascular condition; hyperplasia; kidney disease;
 neovascular condition of the retina; ss.

Homo sapiens.

MO200078341-A1.

28-DEC-2000.

21-JUN-2000; 2000MO-AU00693.

21-JUN-1999; 99US-0140345.

(MURD-) MURDOCH CHILDRENS RES INST.

Wraight CJ, Werther GA, Edmondson SR;

WPI; 2001-041421/05.

Ameliorating the effects of a disorder, e.g. psoriasis, by
 administering UV (ultra-violet) treatment (optional) and an antisense
 nucleic acid that inhibits or reduces growth factor mediated cell
 proliferation and/or inflammation -

Example 7; Page 54; 201pp; English.

The present invention relates to a method for ameliorating the effects
 of skin disorders. The method comprises contacting the skin with an
 antisense oligonucleotide, (for Insulin-like Growth Factor [IGF]-1
 receptor, IGF binding protein [IGFBP]-2 or IGFBP3), which is capable of
 inhibiting or reducing growth factor mediated cell proliferation,
 inflammation and/or other disorders. The present sequence is an
 oligonucleotide which can be used to design the antisense
 oligonucleotides of the present invention (see AAF45151 and
 AAF45153-F45161). The method is useful for ameliorating the effects of
 psoriasis, ichthyosis, pityriasis, ruba, pilaris, serborrhea, keloids,
 keratosis, neoplasias, scleroderma, warts, benign growths, cancers of the
 skin, a hyperneovascular condition such as a neovascular condition of the
 retina, brain or skin, growth factor-mediated malignancies, other
 blood vessels or any other hyperplasia.

Sequence 15 BP; 7 A; 0 C; 3 G; 5 T; 0 other;

Query Match 4.0%; Score 13; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.8e+04; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 agaagaatgttta 38
|||||
Db 2 agaagaatgttta 14

RESULT 43

AAFA8099

ID AAFA8099 standard; DNA; 15 BP.

XX AAFA8099;

XX 30-MAR-2001 (first entry)

DE IGFBP3 oligonucleotide #1519.

XX Antisense therapy; antiproliferative; antiinflammatory; antipsoriatic;
KW cytosolic; dermatological; cardiant; virucide; ophthalmological; keloid;
KW skin disorder; insulin-like Growth Factor 1 receptor; IGF-1; pityriasis;
KW IGF binding protein; IGFBP-2; IGFBP3; inflammation; psoriasis; pilaris;
KW growth factor mediated cell proliferation; ichthyosis; seborrhea; ruba;
KW keratosis; neoplasia; scleroderma; wart; skin cancer; sclerotic disease;
KW hypervascular condition; hyperplasia; kidney disease;
KW neovascular condition of the retina; ss.

XX Homo sapiens.

XX WO200078341-A1.

XX 28-DEC-2000.

XX 21-JUN-2000; 2000WO-AU00693.

XX 21-JUN-1999; 99US-0140345.

XX (MORD-) MORDCH CHILDRENS RES INST.

XX Wraight CJ, Werther GA, Edmondson SR;

XX WPI: 2001-041421/05.

XX Ameliorating the effects of a disorder, e.g. psoriasis, by
PT administering UV (ultra-violet) treatment (optional) and an antisense
PT nucleic acid that inhibits or reduces growth factor mediated cell
PT proliferation and/or inflammation -

XX Example 7; Page 54; 201pp; English.

XX The present invention relates to a method for ameliorating the effects
CC of skin disorders. The method comprises contacting the skin with an
CC antisense oligonucleotide, (for Insulin-like Growth Factor [IGF]-1
CC receptor, IGF binding protein [IGFBP]-2 or IGFBP3), which is capable of
CC inhibiting or reducing growth factor mediated cell proliferation,
CC inflammation and/or other disorders. The present sequence is an
CC oligonucleotide which can be used to design the antisense
CC oligonucleotides of the present invention (see AAF45151 and
CC AAF45153-45161). The method is useful for ameliorating the effects of
CC psoriasis, ichthyosis, pityriasis, ruba, pilaris, seborrhea, keloids,
CC keratosis, neoplasia, scleroderma, warts, benign growths, cancers of the
CC skin, a hypervascular condition such as a neovascular condition of the
CC retina, brain or skin, growth factor-mediated malignancies, other
CC sclerotic disease, kidney disease, hyperproliferation of the inside of
CC blood vessels or any other hyperplasia.

XX Sequence 15 BP; 6 A; 0 C; 4 G; 5 T; 0 other;

Query Match 4.0%; Score 13; DB 22; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.8e+04; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 agaagaatgttta 38
|||||
Db 1 agaagaatgttta 13

RESULT 44

AAA57758/C

ID AAA57758 standard; DNA; 16 BP.

XX AAA57758;

XX 20-OCT-2000 (first entry)

DE Nucleotide sequence which is bound by 22 domain of RIP60 polypeptide.

XX Human; RIP60; zinc finger protein; nucleic acid delivery complex;

XX nucleic acid binding domain; nucleic acid condensation domain; ss.

XX Synthetic.

XX WO200040723-A2.

XX 13-JUL-2000.

XX 04-JAN-2000; 2000WO-US00212.

XX 04-JAN-1999; 99US-0114743.

XX 04-JAN-1999; 99US-0114745.

XX (UYVE-) UNIV VERMONT & STATE AGRIC COLLEGE.

XX Heintz NH, Houchens CR;

XX WPI: 2000-465985/40.

XX Non-viral nucleic acid delivery complex for delivering a nucleic acid
PT molecule into a cell comprises a modular polypeptide -

XX Example 17; Page 74; 115pp; English.

XX AAA57752-66 represent sequences which are bound by the 22 domain of
CC the human RIP60 polypeptide. RIP60 is a zinc finger protein. The
CC nucleic acid binding domain of the RIP60 polypeptide is used to
CC construct a non-viral nucleic acid delivery complex comprising a
CC modular polypeptide. The complex comprises a modular peptide containing
CC a nucleic acid binding domain and a nucleic acid condensation domain
CC that bind with and condense a nucleic acid molecule of more than
CC 50 kilobases in length. The complex also comprises one or more
CC polypeptides selected from a cell recognition domain, a protein
CC transduction domain, a protein degradation domain, an intracellular
CC targeting domain, a protein interaction domain, an epitope domain and
CC a protein purification domain. The complexes are used to deliver a
CC nucleic acid to a cell. The nucleic acids delivered are of various
CC sizes and preferably greater than 50 kilobases, especially more than
CC 100 or more than 200 kilobases in length.

XX Sequence 16 BP; 5 A; 0 C; 0 G; 11 T; 0 other;

Query Match 4.0%; Score 13; DB 21; Length 16;

Best Local Similarity 100.0%; Pred. No. 1.8e+04; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 110 ttaataataataa 122
|||||
Db 15 TTAATAATAATAA 3

RESULT 45

AA092084/c
 ID AA092084 standard; cDNA; 17 bp.
 XX
 AC AA092084;
 XX
 DT 07-JAN-1996 (first entry)
 XX
 DE Renilla reniformis luciferase DNA probe-1.
 XX
 KW Luciferase; enzyme; bioluminescence; luminescence; label; DNA probe;
 KW antibody; oligonucleotide; ss.
 XX
 OS Synthetic.
 XX
 PN US5418155-A.
 PD 23-MAY-1995.
 XX
 PF 29-DEC-1989; 89US-0458952.
 XX
 PR 29-DEC-1989; 89US-0458952.
 PR 20-AUG-1992; 92US-0933017.
 PR 17-JUN-1993; 93US-0079700.
 PR 14-DEC-1993; 93US-0167650.
 XX
 PA (UYGE-) UNIV GEORGIA RES FOUND INC.
 XX
 PI Cormier MJ, Lorenz WW;
 XX
 DR WPI; 1995-199740/26.
 XX
 PT New recombinant Renilla luciferase polypeptide - used as a
 PT luminescent tag, partic in bio-luminescence assays and for the prodn
 PT of antibodies
 XX
 PS Disclosure; Fig. 4; 18bp; English.
 XX
 CC This 17-mer oligonucleotide DNA probe, along with Probe-2 (AA092085)
 CC are used to screen an R. reniformis cDNA library to isolate cDNA
 CC encoding Renilla luciferase. The luciferase was then expressed
 CC using E. coli.
 XX
 SQ Sequence 17 BP; 6 A; 0 C; 2 G; 9 T; 0 other;
 XX

Query Match 4.0%; Score 13; DB 16; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 271 aaaaattattt 283
 |||||||||
 DB 15 AAAAAATTATT 3

Search completed: January 24, 2002, 03:28:19
 Job time: 3671 sec

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Page 1

Oligonucleotide D1
Oligonucleotide D1
Oligonucleotide D2
Oligonucleotide D2
Oligonucleotide D1
Human prothrombin

Page 1
Can be
removed

Page 1

Oligonucleotide D1
Solanium tuberosum

B. burgdorferi ant
B. burgdorferi ant

Human CDNA clone (sequenc

Human genomic DNA
Soybean 318013 reg

Aromatic acyl tran
Human secreted pro

Clone pACCl encodi
Zucchini ACC synth
Zucchini 1-aminocy

Zucchini ACC synth
Plasmodium falcipa

Nucleotide sequenc
Human secreted pro

Human cDNA clone (*Arabidopsis thaliana*)

C	12	30.6	34.0	936	22	AAE58254	Oligonucleotide D1
C	13	30.6	34.0	936	22	AAE58257	Oligonucleotide D1
C	14	30.6	34.0	936	22	AAE58259	Oligonucleotide D2
C	15	30.6	34.0	936	22	AAE58262	Oligonucleotide D2
C	16	30.6	34.0	938	22	AAE58255	Oligonucleotide D1
C	17	30.6	34.0	26928	20	AAZ32184	Human prochromin
C	18	30.6	34.0	87350	18	AAH33003	Human WRN genomic
C	19	30.4	33.8	1347	20	AAXI3338	Enterococcus faecae
C	20	30.4	33.8	2364	21	AAA70246	Plasmidium falcipa
C	21	30	33.3	90	20	AAK60301	Fragment of the be
C	22	30	33.3	244	22	AAE58238	Solanum tuberosum
C	23	30	33.3	4140	18	AAT77331	B. burgdorferi ant
C	24	29.8	33.1	210	20	AAK61492	B. burgdorferi ant
C	25	29.8	33.1	324	20	AAK61491	Human cDNA clone (
C	26	29.8	33.1	784	22	AAH05038	Human cDNA sequenc
C	27	29.8	33.1	3454	22	AAH18466	Human genomic DNA
C	28	29.8	33.1	14417	22	AAI62923	Human genomic DNA
C	29	29.8	33.1	14426	22	AAI62921	Human genomic DNA
C	30	29.8	33.1	513445	22	AAI61373	Soybean 318013 reg
C	31	29.6	32.9	1518	17	AAT67313	Aromatic acyl tran
C	32	29.6	32.9	1592	21	AAZ57085	Human secreted pro
C	33	29.6	32.9	1703	22	AAO51311	Clone PACCl encodi
C	34	29.6	32.9	1703	19	AAV15701	Zucchini ACC synth
C	35	29.6	32.9	1703	22	AAAD0541	Zucchini 1-aminocy
C	36	29.6	32.9	1703	22	AAEF33618	Zucchini ACC synth
C	37	29.6	32.9	5994	21	AAAT0222	Plasmodium falcipa
C	38	29.6	32.9	110000	22	AAAF4800	Nucleotide sequenc
C	39	29.4	32.7	393	21	AACQ9962	Human secreted pro
C	40	29.4	32.7	546	22	AAH09430	Human cDNA clone (
C	41	29.4	32.7	1219	21	AAAC8516	Arabidopsis thalia

ALIGNMENTS

42	32.7	1222	21	AAC45059	Arbidopsis thalia
29.4	32.7	1223	21	AAC00150	Arbidopsis thalia
43	29.4	1223	21	AAC00150	Arbidopsis thalia
44	29.4	32.7	1405	AAH34185	Human colon cancer
45	29.4	32.7	1978	AAH14119	Human cDNA sequenc

Arabidopsis thalia
Arabidopsis thalia
Human colon cancer
Human cDNA sequenc

XX The present sequence encodes the beta-2 toxin of Clostridium
 CC perfringens type C. The specification describes the Clostridium
 CC perfringens beta 2 toxin gene promoter (see AAX60300). The
 CC sequences can be used to produce vaccines against Clostridium,
 CC and especially Clostridium perfringens, or Clostridium
 CC tetani.
 XX
 SQ Sequence 1392 BP; 606 A; 115 C; 209 G; 462 T; 0 other;

Query Match 100.0%; Score 90; DB 20; Length 1392;
 Best Local Similarity 100.0%; Pred. No. 9,9e-13;
 Matches 90; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atgaaaaaatatttcaagaattactgttaattttatgttttcaatgtttcttatgtt 60
 |||
 DB 268 atgaaaaaatatttcaagaattactgttaattttatgttttcaatgtttcttatgtt 327

QY 61 ggagcaataagtcacaaagcaagtcga 90
 |||
 DB 328 ggagcaataagtcacaaagcaagtcga 357

RESULT 2
 AAX60300
 ID AAX60300 standard; DNA; 327 BP.

AC AAX60300;
 XX
 DT 12-AUG-1999 (first entry)

DE Promoter of the beta-2 toxin gene of Clostridium perfringens type C.

XX Beta-2 toxin; Clostridium perfringens type C; gene promoter;
 XX vaccine; Clostridium tetani; ss.

OS Clostridium perfringens.

PN FR2768747-A1.

XX 26-MAR-1999.

PF 19-SEP-1997; 97FR-0011710.

PR 19-SEP-1997; 97FR-0011710.

PA (INSP) INST PASTEUR.

PI Gilbert M, Popoff MR;

XX MPI; 1999-217498/19.

XX Clostridium beta2 toxin gene promoter and signal sequence - useful
 PT against toxins from Clostridium perfringens

PS Claim 1; Page 32; 46pp; French.

CC The present sequence represents the promoter of the beta-2 toxin
 CC gene of Clostridium perfringens type C. The beta2-toxin promoter
 CC and gene sequences can be used to produce vaccines against Clostridium,
 CC and especially Clostridium perfringens, or Clostridium
 CC tetani.

SQ Sequence 327 BP; 141 A; 13 C; 44 G; 129 T; 0 other;

Query Match 66.7%; Score 60; DB 20; Length 327;
 Best Local Similarity 100.0%; Pred. No. 8,7e-06;
 Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atgaaaaaatatttcaagaattactgttaattttatgttttcaatgtttcttatgtt 60
 |||

DB 268 atgaaaaaatatttcaagaattactgttaattttatgttttcaatgtttcttatgtt 327

RESULT 3
 AAX20260/c
 ID AAX20260 standard; DNA; 9542 BP.

AC AAX20260;

DT 04-MAY-1999 (first entry)

DE Borrelia burgdorferi polynucleotide sequence #13.

KW Borrelia burgdorferi; spirochete; bacterium; pathogen; Lyme disease;
 KW epidemic relapsing fever; endemic relapsing fever; Lyme borreliosis;
 KW infection; diagnosis; characterisation; detection; ds.

OS Borrelia burgdorferi.

PN WO9858943-A1.

XX 30-DEC-1998.

PF 18-JUN-1998; 98WO-US12764.

PR 03-SEP-1997; 97US-0057483.

PR 20-JUN-1997; 97US-0050359.

PR 22-JUL-1997; 97US-0053344.

PR 22-JUL-1997; 97US-0053377.

XX (HUMA-) HUMAN GENOME SCI INC.

PA (MEDI-) MEDIMUNE INC.

XX Clayton R, Dougherty BA, Fraser C, Lathigra R, Smith HO;

PI White OR;

XX MPI; 1999-081217/07.

XX New isolated Borrelia burgdorferi nucleic acids - used to develop

PT products for the detection, diagnosis, characterisation, prevention

PT and therapy of infections, particularly Lyme disease

XX Claim 1; Page 920-925; 1128pp; English.

CC AAX20248 to AAX20402 represent polynucleotide sequences isolated from
 CC Borrelia burgdorferi (Bb). Products derived from Bb can be used for
 CC the detection, diagnosis, characterisation, prevention and therapy of
 CC Bb infections, e.g. Lyme disease. They can also be used for the
 CC production of biosynthetic products, e.g. enzymes. Borrelia belongs
 CC to a family of motile, spiral-shaped bacteria called Spirochetes.
 CC Spirochetes are pathogenic in humans and Borrelia causes epidemic and
 CC endemic relapsing fever, and Lyme borreliosis, more commonly known as
 CC Lyme disease.

SQ Sequence 9542 BP; 3812 A; 1160 C; 1113 G; 3457 T; 0 other;

Query Match 36.2%; Score 32.6; DB 20; Length 9542;
 Best Local Similarity 60.9%; Pred. No. 22;
 Matches 53; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

QY 4 aaaaaaatatttcaagaattactgttaattttatgttttcaatgtttcttatgttga 63
 |||

DB 3932 AAAATRAACATAGCTRAATTAATTTTATATTAATTAATTTTCAATATCTGT 3873

QY 64 gcaataagtcacaaagcaagtcga 90
 |||

DB 3872 GAACATATTATTAATAAAGAGAGACGA 3846

RESULT 4
 AAS01490/c
 ID AAS01490 standard; DNA; 4015 BP.

XX	AA01490;
AC	
XX	18-JUL-2001 (first entry)
DT	
XX	
DE	Human secreted protein gene #31.
XX	
KW	Human secreted protein; gene therapy; autoimmune disease;
KW	hyperproliferative disorder; cardiovascular disorder;
KW	cerebrovascular disorder; nervous system disorder; infection;
KW	ocular disorder; wound healing; epithelial cell proliferation;
KW	skin aging; transplantation; tissue regeneration; chemotaxis;
KW	food additive; preservative; ds.
XX	
OS	Homo sapiens.
PN	MO200123402-AI.
PD	05-APR-2001.
PF	26-SEP-2000; 2000WO-US26376.
PR	27-SEP-1999; 99US-0155808.
XX	
XX	(HUMA-) HUMAN GENOME SCI INC.
PA	
PI	Rosen CA, Ruben SM, Komatsoulis GA;
DR	WPI: 2001-266138/27.
DR	P-PsDB: AAU01070.
PT	Nucleic acids encoding 43 human secreted polypeptides, useful for
PT	preventing, diagnosing and/or treating e.g. cancers, Parkinson's
PT	disease and diabetic retinopathy -
PS	Claim 4; Page 452-453; 516pp; English.
XX	
CC	AA01460-AA01502 encode for novel human secreted proteins. The
CC	invention relates to 43 novel human secreted proteins (AAU01040-AAU01082)
CC	and their gene sequences which can be used in gene therapy. The secreted
CC	proteins are useful to prevent, treat or ameliorate a medical condition
CC	in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or
CC	sheep. The secreted proteins are also useful in diagnosing a pathological
CC	condition or susceptibility to a pathological condition. Antibodies to
CC	the secreted proteins can also be used in alleviating symptoms associated
CC	with disorders and in diagnostic immunoassays e.g. radioimmunoassays or
CC	enzyme linked immunosorbent assays (ELISA). Disorders which are diagnosed
CC	or treated include autoimmune diseases e.g. rheumatoid arthritis,
CC	hyperproliferative disorders e.g. neoplasms of the breast or liver,
CC	cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders
CC	e.g. cerebral ischemia, angiogenesis, nervous system disorders e.g.
CC	Parkinson's disease, infections caused by bacteria, viruses and fungi and
CC	ocular disorders e.g. corneal infection. The polypeptides can also be
CC	used to aid wound healing and epithelial cell proliferation, to prevent
CC	skin aging due to sunburn, to maintain organs before transplantation, for
CC	supporting cell culture of primary tissues, to regenerate tissues and in
CC	chemotaxis. The polypeptides can also be used as a food additive or
CC	preservative to increase or decrease storage capabilities.
XX	
SO	Sequence 4015 BP; 1122 A; 712 C; 743 G; 1438 T; 0 other;
<hr/>	
Query Match	36.0%; Score 32.4; DB 22; Length 4015;
Best Local Similarity	68.2%; Pred. No. 24;
Matches	45; Conservative 0; Mismatches 21; Indels 0; Gaps 0
OY	2 tgaaaaaatttcaaaagttaactgtaattttatgtttcattcttatgttg 61
Dd	3088 TGAGGAAGAAGGTATGCAAAAGTTACAGTCATTGTATCTTTCATTTCATGTAAGTAGTG 3029
OY	62 gagcaa 67
Dd	3028 GAGCCTA 3023

```

RESULT      6
TAT34620/c

AAA62306/c
ID AAA62306 standard; DNA; 6250 BP.
XX
AC AAA62306;
XX
DT 12-JAN-2001 (first entry)
XX
DE Caenorhabditis elegans pdk-1 gene.
XX
KW Caenorhabditis elegans; pdk-1b; AKT kinase; daf-18;
KM insulin signalling pathway; daf-2; age-1; insulin receptor; PI 3-kinase;
KV PKB kinase; PTEN lipid phosphatase; antidiabetic; anorectic; obesity;
XX diabetes; ds.
OS
SN Caenorhabditis elegans.
PN WO200033068-A1.
PP 08-JUN-2000.
PR 02-DEC-1999; 99WO-US28529.
PR 03-DEC-1998; 98US-0205658.
PA (GEHO ) GEN HOSPITAL CORP.
PI Ruvkun G, Ogg S;
PI WPI: 2000-423022/36.
DR P-PADB; AAB06177, AAB06178.
PT Diagnosing and treating obesity and impaired glucose tolerance using
XX modulators of daf-18 expression and/or activity -
PS Disclosure: Fig 35; 402pp; English.
XX
XX The present sequence is the genomic sequence of pdk-1 from Caenorhabditis
CC elegans. It encodes two alternatively spliced isoforms which shows
CC homology to mammalian PKC. A number of C. elegans genes have been
CC identified as homologues of genes in the mammalian insulin signalling
CC pathway. The C. elegans age-1 gene encodes a homologue of the mammalian
CC PI 3-kinase whilst daf-2 encodes a homologue of the mammalian insulin
CC receptor. The C. elegans AKT kinase and PKB kinase act downstream of
CC daf-2 and age-1, just as their mammalian homologues act downstream of
CC insulin signalling. The C. elegans PTEN lipid phosphatase homologue,
CC DAF-18, has been found to act upstream of AKT in the pathway. This
CC discovery has enabled mammalian PTEN action to be mapped to the insulin
CC signalling pathway. Conserved DAF motifs can be used to design probes to
CC identify mammalian DAF homologues and thus to identify individuals with
CC a predisposition toward the development of glucose intolerance
CC conditions, such as obesity and diabetes.
XX
SQ Sequence 6250 BP; 1972 A; 1225 C; 1140 G; 1913 T; 0 other;
```

```

ID  AAT34620 standard; DNA; 3337 BP.
XX
XX  AAT34620;
AC
XX  12-NOV-1996 (first entry)
DT
XX  P. vivax ESP-1 blood stage antigen coding sequence.
DE
XX  ESP-1; blood stage antigen; diagnosis; malaria; infection;
KM  causative agent; antibody; monoclonal; polyclonal; assay; ds.
XX
XX  Plasmodium vivax (clone PvMB3.3.1).
OS
XX
XX  Key      Location/Qualifiers
FH  Exon     1..91
FT  Exon     /tag= a
FT  Exon     /note= "encodes initial (N-terminal) sequence of
FT  Exon     /note= "hydrophobic amino acids"
FT  Intron    92..230
FT  Intron    /tag= b
FT  Intron    /note= "contains typical malaria intervening
FT  Intron    /note= "sequence splice sites splice sites"
FT  Exon      231..3197
FT  Exon      /tag= c
FT
FT
XX  US5532133-A.
XX
XX  02-JUL-1996.
XX
XX  02-JUN-1993; 93US-0072610.
XX
XX  02-JUN-1993; 93US-0072610.
XX
XX  (UNIV ) UNIV NEW YORK STATE.
XX
XX  Barnwell JW;
XX
XX  WPI; 1996-32110/32.
XX
XX  P-PSDB; AAR98747.
XX
XX  Antibodies to Plasmodium vivax blood stage antigens - used to
PT  diagnose malaria and to determine whether P. vivax is the species
PT  responsible for infection
XX
XX  Example 4; Column 15-20; 22pp; English.
XX
XX  The present sequence encodes a species-specific Plasmodium vivax
CC  malarial antigen, PvESP-1. The gene appears to be missing a small
CC  portion of its 5' end. This protein is secreted into the plasma of
CC  a susceptible mammalian host after infection. Monoclonal/polyclonal
CC  antibodies can be utilized in assays used to diagnose malaria, as well
CC  as to determine whether P. vivax is the species responsible for the
CC  infection.
XX
XX  Sequence 3337 BP; 1304 A; 467 C; 875 G; 691 T; 0 other;
SQ
Query Match      34.7%; Score 31.2; DB 17; Length 3337;
Best Local Similarity 63.2%; Pred. No. 44;
Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
OY  10 atattcaaaagttactgtatattatgtttcatgtttctattgttgagcaata 69
DB  138 AATATATATAGTATATATAGCAAGTTTATAGATTATTTTATCTTACTGTGAACATGA 79
OY  70 agtccaatgaagaacaa 85
DB  78 ATTAAAGAAAGCAAA 63
RESULT 7
ID AAX15174 standard; DNA; 3337 BP.

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```

XX  AAX15174;
AC
XX  28-APR-1999 (first entry)
DT
XX  DNA encoding a secreted blood-stage protein called PvESP-1.
DE
XX  Erythrocyte secreted protein-1; PvESP-1; malarial antigen;
KM  blood-stage protein; malaria; monoclonal antibody 1D11G10; ds.
XX
XX  Plasmodium vivax.
OS
XX
XX  Key      Location/Qualifiers
FH  CDS      1..3197
FT  CDS      /tag= a
FT  CDS      /note= "contains 1 intron"
FT  Exon      1..91
FT  Exon      /tag= b
FT  Exon      /number= 1
FT  Intron    92..230
FT  Intron    /tag= c
FT  Intron    /number= 1
FT  Exon      231..3194
FT  Exon      /tag= d
FT  Exon      /number= 2
FT
FT
XX  US5874527-A.
XX
XX  23-FEB-1999.
XX
XX  30-SEP-1996; 96US-0719822.
XX
XX  02-JUN-1993; 93US-0072610.
XX
XX  07-JUN-1995; 95US-0478417.
XX
XX  30-SEP-1996; 96US-0719822.
XX
XX  (UNIV ) UNIV NEW YORK STATE.
XX
XX  Barnwell JW;
XX
XX  WPI; 1999-180063/15.
XX
XX  P-PSDB; AAW97039.
XX
XX  Plasmodium vivax peptide antigen - for diagnosis of malaria caused
PT  by Plasmodium vivax
XX
XX  Example 4; Fig 5A-C; 23pp; English.
XX
XX  The present sequence encodes a C-terminal erythrocyte secreted
CC  protein-1 (PvESP-1) of Plasmodium vivax. PvESP-1 is a malarial
CC  antigen which is a secreted blood-stage protein present in detectable
CC  amounts in biological samples from individuals infected with P. vivax.
CC  The protein comprises an epitope not present in other Plasmodium species
CC  that cause malaria in humans, and is bound by monoclonal antibody
CC  1D11G10. The peptide antigen can be used in immunoassays for diagnosis
CC  of malaria caused by P. vivax and/or can be used to produce antibodies
CC  for use in such immunoassays.
XX
XX  Sequence 3337 BP; 1304 A; 467 C; 875 G; 691 T; 0 other;
SQ
Query Match      34.7%; Score 31.2; DB 20; Length 3337;
Best Local Similarity 63.2%; Pred. No. 44;
Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
OY  10 atattcaaaagttactgtatattatgtttcatgtttctattgttgagcaata 69
DB  138 AATATATATAGTATATATAGCAAGTTTATAGATTATTTTATCTTACTGTGAACATGA 79
OY  70 agtccaatgaagaacaa 85
DB  78 ATTAAAGAAAGCAAA 63

```



```
RESULT 8
AAH76457/c
ID AAH76457 standard; DNA; 3337 BP.
XX
AC AAH76457;
XX
DT 22-OCT-2001 (first entry)
XX
DE Plasmodium vivax ESP-1 DNA.
XX
KW Plasmodium vivax: ESP-1; erythrocyte secreted protein-1; PvESP-1;
KW species-specific; malarial peptide antigen; infection; diagnosis;
KW malaria; ds.
XX
OS Plasmodium vivax.
XX
FH Key Location/Qualifiers
FT CDS 1..3197
FT /tag= a
FT /product= "ESP-1"
FT exon 1..91
FT /tag= b
FT /number= 1
FT intron 92..230
FT /tag= c
FT /number= 1
FT exon 231..3197
FT /tag= d
FT /number= 2
XX
XX US6231861-B1.
XX
PD 15-MAY-2001.
XX
PF 05-JUN-1998; 98US-0092458.
XX
PR 02-JUN-1993; 93US-0072610.
PR 07-JUN-1995; 95US-0478417.
PR 30-SEP-1996; 96US-0719822.
XX
PA (UWNY ) UNIV NEW YORK STATE.
XX
PI Barnwell JM;
DR WPI: 2001-335068/35.
DR P-PSDB; AAG66528.
XX
XX New species-specific Plasmodium vivax malarial peptide antigens,
XX proteins or fragments secreted into the plasma of susceptible mammalian
XX host after infection, useful for diagnosing malaria
XX
XX Example 4; Fig 5; 23pp; English.
XX
XX The invention relates to novel species-specific Plasmodium vivax
XX malarial peptide antigens which are proteins or fragments of
XX proteins secreted into the plasma of a susceptible mammalian host after
XX infection, and to monoclonal or polyclonal antibodies directed against
XX those antigens. The peptide antigens, monoclonal antibodies, and/or
XX polyclonal antibodies are useful in assays to diagnose malaria, and to
XX determine which P. vivax species is responsible for the infection.
XX
XX The present sequence encodes P. vivax erythrocyte secreted
XX protein-1 (PvESP-1), a secreted species-specific blood stage
XX antigen provided in the invention.
XX
XX Sequence 3337 BP; 1304 A; 467 C; 875 G; 691 T; 0 other;
XX
Query Match 34.7%; Score 31.2; DB 22; Length 3337;
Best Local Similarity 63.2%; Pred. No. 44;
Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
QY 10 attattcaagttactgtatttattgtttcattgtttctattgttgagcagata 69
```

```
DB 138 AATATATATGTTATATACACGTTTATTAGATTATTATTTACTTGTGAACGATCA 79
QY 70 agtccaatgaagcaaa 85
DB 78 ATTAAAGAAAGCAAA 63
RESULT 9
AAC35351
ID AAC35351 standard; DNA; 1226 BP.
XX
AC AAC35351;
XX
DT 17-OCT-2000 (first entry)
XX
DE Arabidopsis thaliana DNA fragment SEQ ID NO: 9873.
XX
KW Hybridisation assay; genetic mapping; gene expression control;
KW protein identification; signal transduction pathway;
KW metabolic pathway; promoter; termination sequence; ss.
XX
XX Arabidopsis thaliana.
XX
XX EP1033405-A2.
XX
PD 06-SEP-2000.
XX
PF 25-FEB-2000; 2000EP-0301439.
XX
XX 25-FEB-1999; 99US-0121825.
XX 05-MAR-1999; 99US-0123180.
XX 09-MAR-1999; 99US-0123548.
XX 23-MAR-1999; 99US-0125788.
XX 25-MAR-1999; 99US-0126264.
XX 29-MAR-1999; 99US-0126785.
XX 01-APR-1999; 99US-0127462.
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XX 14-JUN-1999; 99US-0139119.
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PR 26-OCT-1999; 990S-0161361.
PR 28-OCT-1999; 990S-0161920.
PR 28-OCT-1999; 990S-0161992.
PR 28-OCT-1999; 990S-0161993.
PR 29-OCT-1999; 990S-0162142.

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Best Local Similarity 70.7%; Pred. No. 53;
Matches 41; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

OY 29 taattttaatgatttcgaatttctattgttggagcaataagttccaatgaagcaag 86.
Db 1157 ttatttttttttcgaatttctgaatgaagcagtaagtaataacagcgag 1214

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RESULT 10
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ID AAD02659 standard; DNA; 13830 BP.
XX
AC AAD02659;
XX
DT 02-MAY-2001 (first entry)
XX
DE Tomato chromosome 5 harbouring the RIN and MC genes.
XX
KW Tomato; RIN; ripening inhibitor; MC; macrocalyx; sepal development;
XX senescence; pathogen infection; ethylene response; transgenic plant; ds.
XX
OS Lycopersicon esculentum.
XX
FH Key
FT misc_signal
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FT /note= "Putative transcription start site of RIN"
FT 1..5694
FT /tag= b
FT /note= "Transcribed region of RIN gene"
FT 211..5489
FT /tag= c
FT /product= "Tomato ripening-inhibitor (RIN) protein"
FT /note= "The specification states that the RIN gene
FT has 9 exons and 8 introns, however the sequence
FT represented in the figure 7 shows 10 exons and
FT 9 introns"
FT 211..395
FT /tag= d
FT /number= "1"
FT 396..3268
FT /tag= e
FT /number= "1"
FT 3269..3347
FT /tag= f
FT /number= "2"
FT 3348..3834
FT /tag= g
FT /number= "2"
FT 3835..3897
FT /tag= h
FT /number= "3"
FT 3898..3919
FT /tag= i
FT /number= "3"
FT 3920..3923
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FT 3924..4149
FT /tag= k
FT /number= "4"
FT 4150..4248
FT /tag= l
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FT 4249..4389
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FT /number= "5"
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FT 4433..4531
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FT exon

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FT 4993..5404
FT /tag= u
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FT /note= "The rin mutation begins at a point within
FT this region"
FT 5405..5489
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FT /note= "MC promoter sequence: This region separates
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FT /tag= y
FT /note= "Transcribed region of MC gene"
FT 8251..13552
FT /tag= z
FT /product= "Tomato macrocalyx (MC) protein"
FT /note= "The coding region has 8 exons and is interrupted
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FT 8440..10594
FT /tag= ab
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FT /note= "The rin mutation terminates within this
FT region of the MC gene"
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FT intron

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PN	WO200104315-A2.	
PD		
PD	18-JAN-2001.	
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PR	12-JUL-1999; 99US-0143364.	
PA	(TEXA) UNIV TEXAS A & M SYSTEM.	
PA	(CORR) CORNELL RES FOUND INC.	
PI	Giovannoni J, Tanksley S, Vrebalov J, Padmanabhan V, Ruezinsky D;	
PI	White R;	
PX		
DR	WPI; 2001-103084/11.	
PX		
PT	New isolated nucleic acid sequence comprising RIN (ripening-inhibitor	
PT	or MC (macrocalyx) genes for use in genetic transformation techniques	
PT	to manipulate a variety of plant characteristics -	
PX		
PS	Disclosure; Fig 7; 167pp; English.	
PX		
CC	The present sequence is tomato chromosome 5 harbouring the RIN and MC	
CC	genes. The invention relates to the RIN (ripening-inhibitor) and MC	
CC	(macrocalyx) genes. The RIN and MC genes are useful in controlling	
CC	of fruit ripening and quality, control of sepal development and	
CC	size, control of senescence, control of pathogen infection, control	
CC	of ethylene response, and DNA markers for assisted breeding. The	
CC	RIN and MC genes are used in genetic transformation techniques to	
CC	manipulate a variety of plant characteristics. Hence these genes	
CC	represent a valuable new tool for the creation of transgenic plants,	
CC	preferably having one or more added beneficial characteristics.	
XQ	Sequence 13630 BP; 4991 A; 1594 C; 1910 G; 5335 T; 0 other;	
PX		

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				Indels	0
				Gaps	0
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Db	7188	ATATAGAAATTTACTACCAATTTTATATCTTCTCATTAATTTCAAAATTTTAAATTTTC	7129		
QY	61	gagagcataagtlccaatgaagaagcaagtgc	90		
Db	7128	TGAATATTTAAACCAATCTAATCTATTCA	7099		

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AC	AAAF58252;
XX	
DT	24-APR-2001 (first entry)
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DE	Oligonucleotide D1835.
XX	
KW	Electron-transfer group; ETM; mismatch; genotyping.
KW	gene expression; ss.
XX	
OS	Synthetic.
XX	
PN	MO200107665-A2.
XX	
PD	01-FEB-2001.

XX 26-JUL-2000; 2000MO-US20476.
PF
XX 26-JUL-1999; 990US-0145695.
PR
PR 17-MAR-2000; 2000US-0190259.
XX
PA (CLIN-) CLINICAL MICRO SENSORS INC.
XX
PI Umek RM;
DR WPI; 2001-159728/16.
XX
XX Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
PT a single surface -
XX
XX
PS Example 6; Page 127; 159pp; English.
XX
XX The present invention relates to a composition comprising two nucleic
CC acids each containing an electron-transfer group (ETM) having
CC different redox potentials. The invention is used for electronic
CC detection of nucleic acids, especially of substitutions (mismatches)
CC and single-nucleotide polymorphisms, e.g. for genotyping,
XX monitoring gene expression.
XX
XX Sequence 936 BP; 4 A; 139 C; 10 G; 7 T; 776 other;

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Oy	1	atgaaataatatttccaagttacgcgttaattttaagtttcgaagtcttcatacgtt	60									
Db	456	www.....	397									
OY	61	ggagcaataagtcacatgaaa	81									
Db	396	TAAAGC.....	376									

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ID	AAFS8254	standard; DNA; 936 BP.
XX		
AC	AAFS8254;	
XX		
DT	24-APR-2001	(first entry)
XX		
DE	Oligonucleotide D1875.	
XX		
KW	Electron-transfer group; ETW; mismatch; genotyping;	
KW	gene expression; ss.	
XX		
OS	Synthetic.	
XX		
PN	WO200107665-A2.	
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PD	01-FEB-2001.	
XX		
PF	26-JUL-2000; 2000WO-US20476.	
XX		
PR	26-JUL-1999; 990US-0145695.	
PR	17-MAR-2000; 2000US-0190259.	
XX		
PA	(CLIN-) CLINICAL MICRO SENSORS INC.	
XX		
PI	Umek RM;	
DR		
WPI:	2001-159728/16.	
XX		
PT	Nucleic acids containing electron-transfer group, useful as labels in	
XX	hybridization assays, e.g. for genotyping, allowing repeat analyses on	

AAF58262/C
 ID AAF58262 standard; DNA; 936 BP.
 XX
 AC AAF58262;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D2007.
 XX
 XX Electron-transfer group; ETM; mismatch; genotyping;
 KM gene expression; ss.
 XX
 OS Synthetic.
 OS
 PN WO200107665-A2.
 XX
 PD 01-FEB-2001.
 XX
 PF 26-JUL-2000; 2000WO-US20476.
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 XX 26-JUL-1999; 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX
 XX (CLIN-) CLINICAL MICRO SENSORS INC.
 PA
 XX
 XX UmeK RM;
 PI
 XX
 DR WPI; 2001-159728/16.
 XX
 PT Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface -
 PS
 XX
 XX Example 6; Page 128; 1599p; English.
 CC
 CC The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 CC
 XX
 SQ Sequence 936 BP; 5 A; 139 C; 10 G; 6 T; 776 other;
 XX

Query Match 34.0%; Score 30.6; DB 22; Length 936;
 Best Local Similarity 3.7%; Pred. No. 59;
 Matches 3; Conservative 62; Mismatches 16; Indels 0; Gaps 0;

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 DB 456 ww 397
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 DB 396 TAAGCWWWWWWWWWWWWWWWW 376

Search completed: January 24, 2002, 02:22:23
 Job time: 1891 sec

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Matches 90; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 1 ATGAAAAAATTATTTCAGAGTTTACTGTAATTTTATGTTTCATGTTTCTTATTGTT 60

QY 61 ggagcaataagtcacatgaagaagtga 90
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DB 61 GGAGCAATAAGTCACATGAAGAAGTGA 90

RESULT 2
LOCUS L77965 1392 bp DNA BCT 28-JUL-1998
DEFINITION Clostridium perfringens C beta 2 toxin gene, complete cds.
ACCESSION L77965
VERSION L77965.1 GI:3342214
KEYWORDS
SOURCE
ORGANISM Clostridium perfringens C.
            Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;
            Clostridium.
REFERENCE 1 (bases 1 to 1392)
AUTHORS Gilbert,M., Jolivet-Reynaud,C. and Popoff,M.R.
TITLE Beta2 toxin, a novel toxin produced by Clostridium perfringens
JOURNAL Gene 203 (1), 65-73 (1997)
MEDLINE 98085977
REMARK Erratum: [[published erratum appears in Gene 1998 Mar
          27;210(1):173]]
          2 (bases 1 to 1392)
REFERENCE Popoff,M.R.
AUTHORS Direct Submission
TITLE Submitted (15-JAN-1998) Toxines Microbiennes, Institut Pasteur,
JOURNAL Paris cedex 15 75724, France
          GSDB:S:76036.
COMMENT [Flatfile retrieved from GSDB Fri Jul 24 15:39:17 1998].
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Matches 90; Conservative 0; Mismatches 0;

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QY 61 ggagcaataagtcacatgaagaagtga 90
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DB 328 GGAGCAATAAGTCACATGAAGAAGTGA 357

RESULT 3
LOCUS AX004613 1392 bp DNA PAT 24-AUG-2000
DEFINITION Sequence 1 from Patent WO9915669.
ACCESSION AX004613
VERSION AX004613.1 GI:9928053
KEYWORDS
SOURCE Clostridium perfringens.
ORGANISM Clostridium perfringens
            Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;
            Clostridium.
REFERENCE 1 (bases 1 to 1392)
AUTHORS Gilbert,M. and Popoff,M.R.
TITLE Clostridium toxin, and method for preparing immunogenic
JOURNAL compositions
          Patent: WO 9915669-A 1 01-APR-1999;
          GIBERT MARYSE (FR); PASTEUR INSTITUT (FR)
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            /db_xref="GI:9928054"
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            NALKNYDINTVNISEDERVNNVBOYREMLDEPKYDPNOOLKSEFILNSOKSDNKEIF
            NVKTEFLNGAIYDMEFTVSSKDKLIVSDMERTKVENGKYLIFSPFTQVCTMDDEL
            AOAIGGVYPTQYSDRFTYADNIIILNFQYATSGSRDLKVEYSVDHMMKDDVKASQ
            MVTGONPDSAKOIRLITKGSFYKIRIRINFTPASIRVGEYGCA"

BASE COUNT      606 a      115 c      209 g      462 t

ORIGIN
Query Match      100.0%; Score 90; DB 6; Length 1392;
Best Local Similarity 100.0%; Pred. No. 1.9e-10;
Matches 90; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atgaaaaaattattcaagttactgtcaattttatgtttcattgttcttattgtt 60
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DB 268 ATGAAAAAATTATTTCAGAGTTTACTGTAATTTTATGTTTCATGTTTCTTATTGTT 327

QY 61 ggagcaataagtcacatgaagaagtga 90
    |||
DB 328 GGAGCAATAAGTCACATGAAGAAGTGA 357

RESULT 4
LOCUS AP003515 54310 bp DNA circular BCT 10-AUG-2001
DEFINITION Clostridium perfringens plasmid pcpl3 DNA, complete sequence.

```


ACCESSION AP003515 GI:15076712
VERSION
KEYWORDS Clostridium perfringens (strain:13) plasmid:pcp13 DNA.
SOURCE Clostridium perfringens
ORGANISM Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae; Clostridium.
REFERENCE 1 (bases 1 to 54310)
AUTHORS Ohnari,K., Ohshima,S., Hirakawa,H., Ohshima,K., Shiba,T., Shimizu,T., Hattori,M., Kuhara,S., Hayashi,H. and Shimizu,T.
TITLE Complete Nucleotide Sequence of the Virulence Plasmid pcp13 from Clostridium perfringens unpublished
JOURNAL 2 (bases 1 to 54310)
REFERENCE Shimizu,T.
AUTHORS
TITLE Direct Submission
JOURNAL Submitted (12-APR-2001) Tohru Shimizu, Institute of Basic Medical Sciences, University of Tsukuba, Department of Microbiology; 1-1-1 Tennohda, Tsukuba, Ibaraki 305-8575, Japan
(E-mail:shimizuemd.tsukuba.ac.jp, Tel:81-298-53-3354, Fax:81-298-53-3354)
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PCP01
para family"
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1751. .3031
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3147. .3509
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/translation="MEKTLAEKRINISPKKNGALVTTLYLPPKMLEVIGITENERE CFEYIEDAKIKSEKQSEFAKEKTIISPKSTFTYLNKKMLEYLGVSSEDRSCITEL RKDITLVKNDGRILDI"
3773. .4024
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3773. .4024
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/note="83 aa, similar to pir:R14710 probable transposase from Yersinia pestis (402 aa); 44% identity in 50 aa overlap
truncated"
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4040. .4222
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complement(5169. .5804)
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/note="211 aa, similar to gp:AP001508.4 ABC transporter (ATP-binding protein) from Bacillus halodurans (213 aa); 49% identity in 214 aa overlap"
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complement(5804. .7966)
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complement(5804. .7966)

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product from Bacillus halodurans (713 aa); 23% identity in
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LEIYQENEDQILKLEIDLESIIYOKRMGSSQYMETLILVQFIDILMIF
YVGLSKYKIGIOKLIGHSTFYMKEKLELVRIYIVLVTVLLVFNKFNPSLF
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ASIIIFPSNADDSISCKGEKRYKWEERKOYIIPELGFNDSIDSPSEEMEK
EBAVYLRKQCAILLADENRIPETSMEEKQMLPEYKRETIIVNPVLKRVLDV
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OKOKIIMKSNQKPSYLLDVNPEGNYVTDIVSVLESNDKLSYKIIQYNSNP
KIRANSEEVINGLEKRYDMVYLIDPYLYLVNASTIINIKAKYKIVIFAIVILLAV
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8779.9012
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9127.9366
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Best Local Similarity 98.9%  Pred. No. 3.9e-10;
Matches 89; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 atgaaaaaattattcaagttactgaatttattgaatttcacggtttcttattgtt 60
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Db 13654 ATGAAAAAATTTTCAAGTTTACTGTAATTTTATGTTTTCATATTTTCTATTGTT 13713

QY 61 ggagcaataagtcacatgaagaagcaatgca 90
|||||
Db 13714 GGAGCAATAAGTCACATGAAGAAGCAACTGCA 13743

RESULT 5
AX004614 327 bp DNA PAT 24-AUG-2000
LOCUS AX004614
DEFINITION Sequence 2 from Patent WO9915669.
ACCESSION AX004614
VERSION AX004614.1 GI:9928055
KEYWORDS

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SOURCE          Clostridium perfringens.
ORGANISM        Clostridium perfringens
Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;
Clostridium.
REFERENCE       1 (bases 1 to 327)
AUTHORS         Gilbert,M. and Popoff,M.R.
TITLE           Clostridium toxin, and method for preparing immunogenic
                compositions
JOURNAL         Patent: WO 9915669-A 2 01-APR-1999;
                GIBERT MARYSE (FR); PASTEUR INSTITUT (FR)
FEATURES        source
                1. 327
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                /db_xref="taxon:1502"
BASE COUNT      141 a 13 c 44 g 129 t
ORIGIN
Query Match      66.7%  Score 60; DB 6; Length 327;
Best Local Similarity 100.0%  Pred. No. 0.00053;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atgaaaaaattattcaagttactgaatttattgaatttcacggtttcttattgtt 60
|||||
Db 268 ATGAAAAAATTTTCAAGTTTACTGTAATTTTATGTTTTCATGTTTCTATTGTT 327

RESULT 6
AC021648/c     128809 bp DNA PRI 01-MAY-2001
LOCUS          Homo sapiens 12 BAC RP11-25114 (Roswell Park Cancer Institute
DEFINITION     Human BAC Library) complete sequence.
ACCESSION      AC021648
VERSION         AC021648.25 GI:13899174
KEYWORDS       HTG.
SOURCE         human.
ORGANISM       Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE       1 (bases 1 to 128809)
AUTHORS         Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-oshan,F.R., Allen,C.,
                Alsdorfs,S.L., Amaratunge,H.C., Are,J.R., Banks,T., Barbara,J.,
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Query Match      42.0% Score 37.8; DB 9; Length 128809;
Best Local Similarity 64.0% Pred. No. 29;
Matches 57; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

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Db 65107 ATGGAATTAAATGTCACAGTCTACTTTAATTATATTCAGATTCATTTAAAT 65048
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OY 61 ggagcaataagttccaatgaaagcaagtgc 89
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Db 65047 GGAAGATGAGAAAAATGAGAAATGTC 65019

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RESULT 7
AC079739 152816 bp DNA HTG 29-SEP-2000
LOCUS Homo sapiens chromosome UNK clone CTD-2012B22, WORKING DRAFT
DEFINITION AC079739
ACCESSION AC079739
VERSION AC079739.2 GI:10305265
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
          1 (bases 1 to 152816)
          The sequence of Homo sapiens clone
          Unpublished
          2 (bases 1 to 152816)
          Waterston,R.H.
REFERENCE JOURNAL
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (10-SEP-2000) Genome Sequencing Center, Washington
          University School of Medicine, 4444 Forest Park Parkway, St. Louis,
          MO 63108, USA
COMMENT On Sep 26, 2000 this sequence version replaced gi:10047917.

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----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_MS2012B22
----- Summary Statistics -----
Sequencing vector: MJ3; 95%
Sequencing vector: plasmid; 5%
Chemistry: Dye-terminator ET; 95% of reads
Chemistry: Dye-terminator Big Dye; 5% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 130608 bases at least Q40
Consensus quality: 137552 bases at least Q30
Consensus quality: 141453 bases at least Q20
Insert size: 150000; agarose-fp
Insert size: 149516; sum-of-ctnigs
Quality coverage: 3.13 in Q20 bases; agarose-fp
Quality coverage: 3.30 in Q20 bases; sum-of-ctnigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 34 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.

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* This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

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* 1156: gap of unknown length
* 1256: contig of 1324 bp in length
* 2580: gap of unknown length
* 2680: contig of 1103 bp in length
* 3783: gap of unknown length
* 3883: contig of 1381 bp in length
* 5264: gap of unknown length
* 5364: gap of 1909 bp in length
* 7272: contig of 1473 bp in length
* 7373: gap of unknown length
* 8846: gap of unknown length
* 8946: contig of 1998 bp in length
* 10944: gap of unknown length
* 11044: contig of 1543 bp in length
* 12587: gap of unknown length
* 12687: contig of 2052 bp in length
* 14739: gap of unknown length
* 14839: contig of 1747 bp in length
* 1586: gap of unknown length
* 1686: gap of 1773 bp in length
* 18459: gap of unknown length
* 18559: contig of 2001 bp in length
* 20560: gap of unknown length
* 20660: contig of 3050 bp in length
* 23710: gap of unknown length
* 23809: contig of 2002 bp in length
* 25810: contig of 2413 bp in length
* 25912: gap of unknown length
* 28324: gap of unknown length
* 28424: contig of 4603 bp in length
* 28425: gap of unknown length
* 3028: gap of unknown length
* 33128: contig of 4107 bp in length
* 37235: gap of unknown length
* 37335: gap of 4555 bp in length
* 41890: gap of unknown length
* 41990: contig of 3258 bp in length
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* 54318: gap of unknown length
* 54417: gap of unknown length
* 54418: contig of 4439 bp in length
* 58856: gap of unknown length
* 58857: gap of 4711 bp in length
* 58957: contig of unknown length
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* 73964: gap of unknown length
* 74064: contig of 4449 bp in length
* 78512: gap of unknown length
* 78513: gap of 6390 bp in length
* 85002: contig of 6390 bp in length
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* 106597: gap of 7974 bp in length
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* 114088: contig of 7391 bp in length
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* 123937: contig of 9749 bp in length
* 124037: gap of unknown length
* 124038: gap of 9825 bp in length
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ORIGIN

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Query Match          40.4%; Score 36.4; DB 2; Length 152816;
Best Local Similarity 64.0%; Pred. No. 57;
Matches 55; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

QY      3  gaaaaaatattcaagaattcactgtaatttttcttcaatgcttcttatgttg 62
          ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 42020  GAAATATATTATTAAATAATCTACTGCTTTTCATATTTTATGTTTTCGTTTTC 42079
          ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY      63  agcaataagtcacgaagaagaagtc 88
          ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 42080  TACCAACAGAACTTATTATGATCAATG 42105

RESULT      8
AC022199/c  165176 bp      DNA      HTG      05-APR-2000
LOCUS      Homo sapiens chromosome 17 clone RP11-145G19 map 17, WORKING DRAFT
DEFINITION
ACCESSION  AC022199
VERSION    AC022199.2 GI:7417783
KEYWORDS   HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
            1 (bases 1 to 165176)
            Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
            Anderson,S., Baldwin,J., Barna,N., Beckery,R., Beda,F.,
            Boguslavsky,L., Boukhgalter,B., Brown,A., Burkett,G., Castle,A.,
            Choepel,Y., Collangelo,M., Collins,S., Collymore,A., Cooke,P.,
            DeArillano,K., Dewar,K., Domino,M., Doyle,M., Fencsator,J.,
            Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J.,
            Gardyna,S., Grant,S., Hagos,B., Heatford,A., Horton,L.,
            Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Kleh,J.,
            Landers,T., Lehoczkzy,J., Levine,R., Lieu,C., Liu,G., Locke,K.,
            MacDonald,P., Margulis,N., McEwan,P., McGurk,A., McKernan,K.,
            McPheeters,R., Meldrum,J., Menues,L., Morrow,J., Naylor,J.,
            Norman,C.H., O'Connor,T., O'Donnell,P., Olivari,T.M., Peterson,K.,
            Pierre,N., Pisani,C., Pollara,V., Raymond,C., Riley,R., Rothman,D.,
            Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,
            Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
            Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
            Zimmer,A. and Zody,M.
            Direct Submission
            Submitted (26-JAN-2000) Whitehead Institute/MIT Center for Genome
            Research, 320 Charles Street, Cambridge, MA 02141, USA
            On Apr 5, 2000 this sequence version replaced gi:6759152.
            All repeats were identified using RepeatMasker:
            Smit, A.F.A. & Green, P. (1996-1997)
            http://ftp.genome.washington.edu/RM/RepeatMasker.html
            ----- Genome Center
            Center: Whitehead Institute/ MIT Center for Genome Research
            Center code: WIBR
            Web site: http://www-seq.wi.mit.edu
            Contact: sequence_submissions@genome.wi.mit.edu
            ----- Project Information
            -----
            Center project name: L5174
            Center clone name: L45C19
            ----- Summary Statistics
            Sequencing vector: M13; M7815; 100% of reads
            Chemistry: Dye-terminator Big Dye; 100% of reads
            Assembly program: Phrap; version 0.960731
            Consensus quality: 157408 bases at least Q40
            Consensus quality: 161068 bases at least Q30
            Consensus quality: 162457 bases at least Q20
            Insert size: 165000; agarose-fp
            Insert size: 164076; sum-of-contrigs

```

Quality coverage: 4.3 in Q20 bases; agarose-1p
Quality coverage: 4.4 in Q20 bases; sum-of-coverage

NOTE: This is a 'working draft' sequence. It currently consists of 12 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 3498: contig of 3498 bp in length
3499 3598: gap of 100 bp
3599 6363: contig of 2765 bp in length
6364 6463: gap of 100 bp
6464 8738: contig of 2275 bp in length
8739 8838: gap of 100 bp
8839 12799: contig of 3961 bp in length
12800 12899: gap of 100 bp
12900 19411: contig of 6512 bp in length
19412 19511: gap of 100 bp
19512 29328: contig of 9717 bp in length
29329 40493: gap of 100 bp
40494 40593: contig of 11165 bp in length
40594 52209: gap of 100 bp
52210 52309: gap of 100 bp
52310 71806: contig of 19497 bp in length
71807 71906: gap of 100 bp
71907 98472: contig of 26566 bp in length
98473 98572: gap of 100 bp
98573 128201: contig of 29629 bp in length
128202 128301: gap of 100 bp
128302 165176: contig of 36875 bp in length

FEATURES

source
1. 165176
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="17"
/map="17"
/clone="RP11-145G19"
/clone_1lb="RPC1-11 Human Male BAC"
1. 3498
/note="assembly-fragment"
3599. 6363
/note="assembly-fragment"
6464. 8738
/note="assembly-fragment"
clone_end:sp6
vector_side:left"
8839. 12799
/note="assembly-fragment"
12900. 19411
/note="assembly-fragment"
19512. 29228
/note="assembly-fragment"
29329. 40493
/note="assembly-fragment"
40594. 52209
/note="assembly-fragment"
52310. 71806
/note="assembly-fragment"
71907. 98472
/note="assembly-fragment"
98573. 128201
/note="assembly-fragment"
128302. 165176
/note="assembly-fragment"
clone_end:17
vector_side:left"
51832 a 30819 c 31782 g 49641 t 1102 others

Query Match 40.4% Score 36.4; DB 2; Length 165176;
Best Local Similarity 64.0%; Pred. No. 57;
Matches 55; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

Oy 3 gaataaatttcaagaattcactgtaattttatgtttcactgattttcattgttg 62
Db 18382 GAAATAATTTTAAATATATACGCTTTTCATATATTTTATGCTTTTTC 18323
Oy 63 agcaataagtcgaatgaagaagcagtg 88
Db 18322 TACCACAGAAATCTTATGTCATG 18297

RESULT 9
AC016788/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 167322)
Britten, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, M.,
Baldwin, J., Barne, N., Beckery, R., Boguslavsky, L., Boukhalter, B.,
Brown, A., Castle, A., Colangelo, M., Collins, S., Collymore, A.,
Cooke, P., Dearrellano, K., Dewar, K., Domino, M., Doneelan, L., Doyle, M.,
Ferreira, P., FitzHugh, W., Forrest, C., Funke, R., Gage, D.,
Galagan, J., Gardyna, S., Grant, G., Hagos, B., Heaford, A., Horton, L.,
Howard, J. C., Johnson, R., Jones, C., Kann, L., Karatas, A., Klein, J.,
Lehoczy, J., Lieu, C., Locke, K., Macdonald, P., Marquis, N.,
McKernan, P., McGurt, A., McKernan, K., McLaughlin, J., Meldrum, J.,
Morrow, J., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P.,
Peterson, K., Pollara, V., Riley, R., Roy, A., Santos, R., Severy, P.,
Stange-Thomann, N., Stojanovic, N., Sudranian, A., Talamas, J.,
Testafaye, S., Tirrell, A., Vassiliev, H., Vo, A., Wheeler, J., Wu, X.,
Wyman, D., Ye, W. J., Zimmer, A. and Zody, M.

TITLE
JOURNAL
COMMENT

Submitted (05-DEC-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 20, 2000 this sequence version replaced g1.9123832.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
Project Information
Center project name: 11.E.12
Center clone name: 11.E.12

Summary Statistics
Sequencing vector: M13; M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 159178 bases at least Q40
Consensus quality: 163354 bases at least Q30
Consensus quality: 164761 bases at least Q20
Insert size: 170000; agarose-1p
Insert size: 165922; sum-of-coverage
Quality coverage: 4.9 in Q20 bases; agarose-1p
Quality coverage: 5.0 in Q20 bases; sum-of-coverage

NOTE: This is a 'working draft' sequence. It currently

misc_feature	/note="assembly_fragment clone end:Sp6 vector.size:left" 4428. .6881
misc_feature	/note="assembly_fragment 6982. .9727
misc_feature	/note="assembly_fragment 9828. .12174
misc_feature	/note="assembly_fragment 12575. .14864
misc_feature	/note="assembly_fragment 14965. .18876
misc_feature	/note="assembly_fragment 18977. .60052
misc_feature	/note="assembly_fragment 60153. .69381
misc_feature	/note="assembly_fragment 69482. .77833
misc_feature	/note="assembly_fragment 77934. .86115
misc_feature	/note="assembly_fragment 86216. .98453
misc_feature	/note="assembly_fragment 98354. .111638
misc_feature	/note="assembly_fragment 116459. .134093
misc_feature	/note="assembly_fragment 134194. .162917
misc_feature	/note="assembly_fragment" 163018. .167322

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/note="assembly_fragment  
clone_end:77  
vector_side:right"  
50484 a 31742 c 33143 g 50550 t 1403 others  
  
tch          40.4%   Score 36.4; DB 2; Length 167322;  
lthal Similarity 64.0%; Pred. No. 57;  
55; Conservative 0; Mismatches 31; Indels 0; Gaps 0  
  
agcaataagtcacatgaaagaacgtg 88  
| | | |  
| | | |  
| | | |  
TTACCCAGAGAATCTTATTAGTCATG 15867  
  
AP003462    189771 bp      DNA              HTG           30-MAR-2001  
Homo sapiens chromosome 11 clone RP11-480C22 map 11q, WORKING DRAFT  
SEQUENCE, 13 unordered pieces.  
AP003462  
AP003462.1 GI:13488923  
HTGS: PHASE1: HTGS: DRAFT.  
Homo sapiens DNA, clone:RP11-480C22.  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.  
1 (bases 1 to 189771)  
Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Yada,T., Tochio,Y., Watanabe,H. and Sakaki,Y.  
Direct Submission  
Submitted (29-MAR-2001) Masahira Hattori, The Institute of Physical  
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);  
1-7-22 Shohiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
Tel:81-45-503-9111, Fax:81-45-503-9170)  
  
----- Genome Center -----  
Center: RIKEN Genomic Sciences Center(GSC)  
Center code: RIKEN  
Web site: http://hgp.gsc.riken.go.jp/  
Contact: hattori@gscc.riken.go.jp  
Project Information  
Center project name: HumDraft11  
Center clone name: RP11-480C22  
Summary Statistics  
Sequencing vector: PCR products; 100% of reads  
Chemistry: Dye-terminator EF-amersham; 100% of reads  
Assembly program: Phrap; version 0.990329  
Consensus quality: 187154 bases at least Q40  
Consensus quality: 188004 bases at least Q30  
Consensus quality: 188284 bases at least Q20  
Insert size: 188571; sum-of-contigs  
Quality coverage: 8.56x in Q20 bases; sum-of-contigs  
-----  
NOTE: This is a 'working draft' sequence. It currently consists of  
13 contigs. The true order of the pieces is not known and the  
order in this sequence record is arbitrary. Gaps between the  
contigs are represented as runs N, but the exact sizes of the gaps  
are unknown. This record will be updated with the finished sequence  
as soon as it is available and the accession number will be  
preserved  
1  
43267 43166 contig of 43166 bp in length  
62804 62703 contig of 19437 bp in length  
84577 84476 contig of 21673 bp in length  
106769 106668 contig of 22092 bp in length  
126420 126319 contig of 19551 bp in length  
141582 141481 contig of 15062 bp in length  
153916 contig of 12335 bp in length
```

```

154017 163781 contig of 9765 bp in length
163882 170760 contig of 6879 bp in length
170861 176864 contig of 6004 bp in length
176965 183161 contig of 6197 bp in length
183262 185847 contig of 2586 bp in length
185948 189771 contig of 3824 bp in length.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 13 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
1 43166: contig of 43166 bp in length
* 43167 43266: gap of 100 bp
* 43267 62703: contig of 19437 bp in length
* 62704 62803: gap of 100 bp
* 62804 84476: contig of 21673 bp in length
* 84477 84576: gap of 100 bp
* 84577 106668: contig of 22092 bp in length
* 106669 106768: gap of 100 bp
* 106769 126319: contig of 19551 bp in length
* 126320 126419: gap of 100 bp
* 126420 141481: contig of 15062 bp in length
* 141482 141581: gap of 100 bp
* 141582 153916: contig of 12335 bp in length
* 153917 154016: gap of 100 bp
* 154017 163781: contig of 9765 bp in length
* 163782 163881: gap of 100 bp
* 163882 170760: contig of 6879 bp in length
* 170761 170860: gap of 100 bp
* 170861 176864: contig of 6004 bp in length
* 176865 176964: gap of 100 bp
* 176965 183161: contig of 6197 bp in length
* 183162 183261: gap of 100 bp
* 183262 185847: contig of 2586 bp in length
* 185848 185947: gap of 100 bp
* 185948 189771: contig of 3824 bp in length.

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FEATURES

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source
1. 189771
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   /db_xref="taxon:9606"
   /chromosome="11"
   /map="11g"
   /clone="RP11-480C22"
1. 43166
   /note="assembly_fragment"
43267..62703
   /note="assembly_fragment"
62804..84476
   /note="assembly_fragment_clone_end:17 vector_side:left"
84577..106668
   /note="assembly_fragment"
106769..126319
   /note="assembly_fragment"
126420..141481
   /note="assembly_fragment"
141582..153916
   /note="assembly_fragment"
154017..163781
   /note="assembly_fragment"
163882..170760
   /note="assembly_fragment"
170861..176864
   /note="assembly_fragment_clone_end:SP6 vector_side:left"
176965..183161
   /note="assembly_fragment"
183262..185847
   /note="assembly_fragment"
185948..189771
   /note="assembly_fragment"
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56729 a 37254 c 36581 g 58007 t 1200 others

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ORIGIN

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Query Match 40.4%; Score 36.4; DB 2; Length 189771;
Best Local Similarity 64.0%; Pred. No. 57;
Matches 55; Conservative 0; Mismatches 31; Indels 0; Gaps 0;
OY 3 gaataaatttttaagttacgtatattatglttcacgtttctatgttg 62
Db 140789 GAAAATATTATTAATAATATTACGCTTTCTATATTTATGTTTCTGCTTTTC 140730
OY 63 agcaataatccaatgaaagcaagtg 88
Db 140729 TACCAACAGCAATCTTATAGTCAATG 140704

RESULT 11
AC074180/C DNA 17-AUG-2000
LOCUS
DEFINITION
AC074180.2 GI:9638218
VERSION
AC074180.2 GI:9638218
KEYWORDS
HTG; HTGS-PHASE1; HTGS-DRAFT.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 193415)
Waterston,R.H.
TITLE
The sequence of Homo sapiens clone
JOURNAL
Unpublished
2 (bases 1 to 193415)
Waterston,R.H.
REFERENCE
Direct Submission
Submitted (15-JUL-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Aug 17, 2000 this sequence version replaced gi.9211440.
COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H.NH0480C22
----- Summary Statistics -----
Sequencing vector: M13, 2%
Sequencing vector: plasmid, 0%
Chemistry: Dye-primer ET; 2% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 187074 bases at least Q40
Consensus quality: 192835 bases at least Q30
Consensus quality: 195619 bases at least Q20
Insert size: 182000; agarose-fp
Insert size: 195112; sum-of-contigs
Quality coverage: 7.73 in Q20 bases; agarose-fp
Quality coverage: 7.34 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 10 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
1 3180: contig of 3180 bp in length
* 3181 3280: gap of unknown length
* 3281 12431: contig of 9151 bp in length
* 12432 12531: gap of unknown length
* 12532 23140: contig of 10609 bp in length

```


23141 23240: gap of unknown length
* 23241 34935: contig of 11695 bp in length
* 34936 35035: gap of unknown length
* 35036 50440: contig of 15405 bp in length
* 50441 50540: gap of unknown length
* 50541 71299: contig of 20759 bp in length
* 71300 71399: gap of unknown length
* 71400 96045: contig of 24646 bp in length
* 96046 96145: gap of unknown length
* 96146 124103: contig of 27958 bp in length
* 124104 124203: gap of unknown length
* 124204 160061: contig of 35858 bp in length
* 160062 160161: gap of unknown length
* 160162 193415: contig of 33254 bp in length.
Location/Qualifiers
1. 193415
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/db_xref="taxon:9606"
/chromosome="11"
/clone="RP11-480C22"
1. 3180
misc_feature
/note="assembly_name:Contig21"
3281. 12431
/note="assembly_name:Contig22"
12532. 23140
misc_feature
/note="assembly_name:Contig23"
23241. 34935
misc_feature
/note="assembly_name:Contig24"
35036. 50440
misc_feature
/note="assembly_name:Contig25"
50541. 71299
misc_feature
/note="assembly_name:Contig26"
71400. 96045
misc_feature
/note="assembly_name:Contig27"
96146. 124103
misc_feature
/note="assembly_name:Contig28"
124204. 160061
misc_feature
/note="assembly_name:Contig29"
160162. 193415
/note="assembly_name:Contig30
clone_end:SP6
vector_side:right"

BASE COUNT 57779 a 38131 c 37181 g 59416 t 908 others
ORIGIN

Query Match 40.4%; Score 36.4; DB 2; Length 193415;
Best Local Similarity 64.0%; Pred. No. 57;
Matches 55; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

QY 3 gaaataattattcaagaattactgtatatttatgtttcatgtttcttattgttg 62
DB 21903 GAAATAATTATTAAATATTACTGCTTTTCATATTATTTTATGTTTCTGTTTTTC 21844
QY 63 agcataagttccatgaagaagcagtg 88
DB 21843 TACCAAGAACTTTATAGTCAATG 21818

RESULT 12
LOCUS CNS01BGA 148 bp mRNA PLN 02-SEP-1999
DEFINITION Botrytis cinerea strain 74 cDNA library under conditions of
nitrogen deprivation.
ACCESSION AL114194
VERSION AL114194.1 GI:5828813
KEYWORDS cDNA library; nitrogen deprivation.
SOURCE Botryotinia fuckeliana
ORGANISM Botryotinia fuckeliana
Eukaryota; Fungi; Ascomycota; Pezizomycotina; Leotiomycetes;
Helotiales; Sclerotiniaceae; Botryotinia.
REFERENCE 1 (bases 1 to 148)
Bliton,F., levis,C., Fortini,D., Pradler,J.M. and Brygoo,Y.

TITLE Direct Submission
JOURNAL Submitted (01-SEP-1999) Phytopathologie, INRA, route de St Cyr,
78026 Versailles, France
REFERENCE 2 (bases 1 to 148)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (01-SEP-1999) Genoscope - Centre National de Sequencage :
CP 5706 91057 Evry cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
COMMENT The cDNA library to be analyzed within the framework of this
project was created using a botrytis cinerea strain which was grown
under conditions of nitrogen deprivation, which is the normal
situation for B. cinerea during its development on its host plant.
The library was produced in an oriented direction, in the pBSII
vector.
Location/Qualifiers
1. 148
/organism="Botryotinia fuckeliana"
/strain="74"
/db_xref="taxon:40559"
/note="Genoscope sequence ID : W52A031"
BASE COUNT 33 a 17 c 25 g 73 t
ORIGIN

Query Match 39.38; Score 35.4; DB 8; Length 148;
Best Local Similarity 66.2%; Pred. No. 1e+02;
Matches 51; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

QY 11 ttattcaagttactgtaattttatgtttcatgtttcttattgttgagcaataa 70
DB 45 TTATTTTTCATTTTCATTTATTTTTCCTTTTTCCTTTTTCCTTTTTCGACCAATTA 104
QY 71 gtccaatgaagaagcagtg 87
DB 105 GAGCATCAACATTTAAGT 121

RESULT 13
AF099924/c 33460 bp DNA INV 09-AUG-2001
LOCUS Caenorhabditis elegans cosmid K07A9, complete sequence.
DEFINITION AF099924
ACCESSION AF099924.2 GI:6671811
VERSION HMG.
KEYWORDS Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;
Rhabditidae; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 33460)
The C. elegans Sequencing Consortium.
JOURNAL Genome sequence of the nematode C. elegans: a platform for
investigating biology. The C. elegans Sequencing Consortium
Science. 282 (5396), 2012-2018 (1998)
99069613
REFERENCE 2 (bases 1 to 33460)
Davidson, S. and O'Neal, D.
TITLE The sequence of C. elegans cosmid K07A9
JOURNAL Unpublished
AUTHORS 3 (bases 1 to 33460)
JOURNAL Direct Submission
TITLE Unpublished
AUTHORS 4 (bases 1 to 33460)
JOURNAL Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (20-OCT-1998) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
5 (bases 1 to 33460)
JOURNAL Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (27-MAY-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

SOURCE ORGANISM human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 132319)
AUTHORS Sulston, J.E. and Waterston, R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792

REFERENCE 2 (bases 1 to 132319)
AUTHORS Ryan, E., Bauer, C., Tuccil, S. and Spalding, L.
TITLE The sequence of Homo sapiens BAC Clone GSI-146J4
JOURNAL Unpublished
3 (bases 1 to 132319)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (11-JAN-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
4 (bases 1 to 132319)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (05-MAY-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
5 (bases 1 to 132319)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (11-SEP-1999) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
6 (bases 1 to 132319)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (21-DEC-1999) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On May 5, 1999 this sequence version replaced g1:4337266.

REFERENCE 6
AUTHORS Center: Washington University Genome Sequencing Center
TITLE Center code: WUGSC
JOURNAL Web site: http://genome.wustl.edu/gsc
Contact: sapiens@wustl.wustl.edu
Summary Statistics
Center project name: H_GSI46J04

COMMENT

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
The sequence of this clone was established as part of a mapping and sequencing collaboration between the NHGRI Chromosome 7 Mapping Project (Eric D. Green, Director), John D. McPherson in the Department of Genetics (Washington University), and the Washington University Genome Sequencing Center. For additional information about the map position of this sequence, see
http://www.nhgri.nih.gov/DIR/CTB/CHR7 send
mailto:egreen@nhgri.nih.gov, or see http://genome.wustl.edu/gsc

SOURCE INFORMATION:
This clone is from the first BAC library from Genome Systems, Inc.
(http://www.genomesystems.com).
Cell line: lymphoblastoid
Haplotypes: two

VECTOR: pBelBAC
Selection: chloramphenicol
NEIGHBORING SEQUENCE INFORMATION:
The actual start of this clone is at base position 1 of GSI-146J4.
Actual end is at 132319 of GSI-146J4.

FEATURES
source
Location/Qualifiers
1..132319
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="7"
/map="7q31.1-31.3"
/clone="GSI-146J4"
/clone_id="GSBAC1"
522..601
/rpt_family="L2"
1216..1506
/rpt_family="Alu"
1514..1858
/rpt_family="L1"
2118..2474
/rpt_family="MALR"
2512..2783
/rpt_family="MER4-group?"
3328..3362
/rpt_family="(TGA)n"
3810..3833
/rpt_family="POLY_G"
3880..4030
/rpt_family="MER1_type"
4108..4397
/rpt_family="Retroviral"
6413..6446
/rpt_family="(CA)n"
7359..7395
/rpt_family="AT-rich"
7396..7704
/rpt_family="MER1_type"
7930..7982
/rpt_family="Retroviral"
7983..8089
/rpt_family="(TA)n"
8111..8253
/rpt_family="Retroviral"
10366..10769
/note="match to EST R59425 (NID:g830120) yhi7c08.r1"
10401..10850
/note="match to EST H18852 (NID:g885092) ym45f11.r1"
10734..11029
/note="match to EST AA328782 (NID:g1981047)"
11459..11522
/rpt_family="L2"
11660..12114
/note="similar to EST A1220397 (NID:g3802600) gq73h11.x1"
11665..11725
/note="similar to Mus musculus EST AA986059 (NID:g3167448) uc72e08.y1"
11713..44221
/gene="WUGSC:H_GSI46J04.1"
1019111713..11725,15668,15772,22543,22572,23259,23382,30494,30552,31735,31785,34555,34639,43413,43539,44132,44221
/gene="WUGSC:H_GSI46J04.1"
/note="match to G92520 (PID:g3334194): H_GSI46J04.1"
/codon_start=1
/product="GS3786"
/protein_id="AAB54511.1"
/db_xref="GI:3670323"
/translation="MRVAGAAKLVAVAVFLLTFYISOVEIRKQDASLGNLPAARSALDTAASTKPPRRYKQISKACPEKHEAFKASGAANVVPKICLEDNVLMGSKNVGRGINVALANGTGEVDTKTFDMGCDVAFIEFLKAIODGTLVLMGTVDGATKLNDEARRLIADIGSTITNIGFRDMNVPCGGKIKTKSPEDGIIKNNKDTNRYEGMPEVEMEGCIPKOD"

misc_feature
11766..12114

Db 105826 GAATAATATTATTTCAGGTCGAATGTATATTTATGCTTAATGTTTTCTTA 1051/4

Search completed: January 24, 2002, 02:27:00
Job time: 4473 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:51:28 ; Search time 2272.52 Seconds
(without alignments)
425.572 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90

Sequence: 1 atgaaaaaatattattcaaa.....gtccaatgaagaagtgcga 90

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 11351937 seqs, 5372889281 residues

Total number of hits satisfying chosen parameters: 22703874

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

EST:*
1: em_estfun:*
2: em_esthum:*
3: em_estin:*
4: em_estom:*
5: em_estpl:*
6: em_estba:*
7: em_estro:*
8: em_estoy:*
9: em_hic:*
10: gb_estl:*
11: gb_est2:*
12: gb_hic:*
13: gb_gss:*
14: em_gss_fun:*
15: em_gss_hum:*
16: em_gss_inv:*
17: em_gss_pln:*
18: em_gss_pro:*
19: em_gss_rod:*
20: em_gss_vrt:*
21: em_gss_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
C 1	37.8	42.0	561	13	AO533830	RPC1-11-3
C 2	35.6	39.6	302	10	AV247937	AV247937
C 3	34.8	38.7	447	13	B67487	T25W7TR TAM
C 4	34.8	38.7	482	13	AO968516	AO968516 LERJDO3TR
C 5	34.8	38.7	488	13	AO968397	AO968397 LERJDO3TR
C 6	34.8	38.7	497	13	AO958266	AO958266 LERJDO3TR
C 7	34.8	38.7	498	13	B77731	T28N11TR TA
C 8	34.8	38.7	623	13	B30114	T26F22TRB T
C 9	34.8	38.7	627	13	B78178	T31J11TR TA
C 10	34.8	38.7	640	13	AO958844	LERJDO3TR
C 11	34.8	38.7	643	13	B98363	T24C11TRB T
C 12	34.8	38.7	660	13	CNS00WZN	AL094241 Arabidops

13	34.8	38.7	683	13	AO968517	AO968517 LERJDO3TR
C 14	34.8	38.7	743	13	AO958265	AO958265 LERJDO3TR
C 15	34.8	38.7	798	13	AO956484	AO956484 LERJDO3TR
C 16	34.4	38.2	607	13	A2522516	A2522516 207PDC02
C 17	34.4	38.2	638	13	A2523727	A2523727 222PBH02
C 18	34.2	38.0	923	13	BH132631	BH132631 ENT07OTR
C 19	34.2	37.8	871	13	A2680048	A2680048 ENT07OTR
C 20	33.8	37.6	867	13	CNS00CX5	AL060052 DROSOPH11
C 21	33.6	37.3	240	10	AU072517	AU072517
C 22	33.6	37.3	240	10	AU072714	AU072714
C 23	33.6	37.3	306	11	C24362	C24362
C 24	33.6	37.3	376	10	AV760889	AV760889
C 25	33.6	37.3	413	13	AO798320	HS_3160_A
C 26	33.6	37.3	552	13	A2065684	A2065684
C 27	33.6	37.3	676	13	AO635998	RPC1-11-4
C 28	33.6	37.3	913	13	BH132369	BH132369
C 29	33.4	37.1	432	10	AA149033	AA149033
C 30	33.2	36.9	463	11	BG603509	BG603509
C 31	33.2	36.9	526	11	BG603424	EST502514
C 32	33.2	36.9	652	11	BG603423	EST502513
C 33	33.2	36.7	1101	13	CNS00K05	AL070967 DROSOPH11
C 34	32.8	36.4	420	11	Z38090	Z38090
C 35	32.8	36.4	473	13	AO001587	AO001587
C 36	32.8	36.4	589	13	AO026620	CIT-HSP-2
C 37	32.8	36.4	843	13	A2549410	ENT07OTR
C 38	32.8	36.4	899	13	BH136018	BH136018
C 39	32.8	36.4	944	13	A2533633	ENT07OTR
C 40	32.8	36.4	949	13	CNS01BD5	AL109198 DROSOPH11
C 41	32.8	36.4	958	13	A2687864	ENT07OTR
C 42	32.6	36.2	510	13	A2384026	IM0141G19
C 43	32.6	36.2	553	13	AO698845	HS_5557_B
C 44	32.4	36.0	366	13	BH014409	TOGBLA2TH
C 45	32.4	36.0	589	13	AZ247746	RPC1-23-9

ALIGNMENTS

RESULT 1
AO533830/c 561 bp DNA 18-MAY-1999
LOCUS
DEFINITION
RPC1-11-379L24.TV RPC1-11 Homo sapiens genomic clone RPC1-11-379L24
ACCESSION
AO533830
VERSION
AO533830.1 GI:4845520
KEYWORDS
GSS.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
Zhao,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and Venter
J.C.
TITLE
Use of BAC End Sequences from Library RPC1-11 for Sequence-Ready
Map Building
JOURNAL
Unpublished (1997)
COMMENT
Other-GSSs: RPC1-11-379L24.TV
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org

Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Pieter de Jong (pieterdejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.
Seq primer: T7
Class: BAC ends.
Location/Qualifiers

FEATURES

Query Match	Best Local Similarity	Match 48: Conservative	Score 73.8%	DB 13: Indels	Length 561: Gaps 0
4	aaaaaatattatccaagttactctgaatttaattgaatttcacgtttcttattgttga	63			
Db	428 AAGAGCAATATTATTCAGTGTCTTGCTGCTTTTTCATTTTCATTTCTTTTATGGTTGA	369			
Qy	64 gcaat 68				
Db	368 GAAAT 364				
RESULT 2	AV247937	302 bp	EST	04-NOV-1999	
LOCUS	AV247937				
DEFINITION	musculus full-length enriched, 0 day neonate head Mus				
ACCESSION	AV247937				
VERSION	AV247937.1	GI:6235396			
KEYWORDS	EST.				
SOURCE	house mouse.				
ORGANISM	Mus musculus				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.				
AUTHORS	1 (bases 1 to 302)				
	Konno,H., Aizawa,K., Akahira,S., Akiyama,J., Carninci,P., Endo,T., Fukuda,S., Fukunishi,Y., Hara,A., Hayatsu,N., Hirazane,T., Hori,F., Ishii,Y., Ishikawa,T., Itoh,M., Izawa,M., Kadota,K., Kagawa,I., Ka,C., Kawai,T., Kikuchi,N., Kojima,Y., Koya,S., Kusakabe,M., Matsuyama,T., Miki,R., Mizuno,Y., Nakamura,M., Oda,H., Okazaki,Y., Owa,C., Ozawa,Y., Saito,H., Sano,M., Sato,K., Shibata,K., Shibata,Y., Shigemoto,Y., Shiraki,T., Sogabe,Y., Sugahara,Y., Suzuki,H., Suzuki,H., Takahashi,F., Tateo,M., Tominaga,N., Tsunoda,Y., Watanishi,A., Watanabe,S., Yamamura,T., Yasunishi,A., Yokota,T., Yoshiki,A., Yoshino,M., Muramatsu,M. and Hayashizaki,Y.				
TITLE	RIKEN Mouse ESTs (Konno,H., et al. 1999)				
COMMENT	Unpublished (1999)				
	Contact: Yoshihide Hayashizaki				
	Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute				
	The Institute of Physical and Chemical Research (RIKEN)				
	1-7-22 Suenho-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan				
	Tel: 81-45-503-9222				
	Fax: 81-45-503-9216				
	Email: genome-res@sc.riken.go.jp,				
	URL: http://genome.gsc.riken.go.jp/				
	Sasaki,N., Iwawa,M., Watanishi,M., Ozawa,K., Tanaka,T., Yoneda,Y., Matsura,S., Carninci,P., Muramatsu,M., Okazaki,Y. and Hayashizaki,Y.				
	Transcriptional sequencing: A method for DNA sequencing using RNA polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)				
	Itoh,M., Kikuchi,T., Akiyama,Y., Shibata,K., Izawa,M., Kawai,J., Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M., Okazaki,Y. and Hayashizaki,Y.				
	Automated filtration-based high-throughput plasmid preparation system. Genome Res. 9 (5), 463-470 (1999)				
	Carninci,P. and Hayashizaki,Y.				

High-efficiency full-length cDNA cloning. *Methods Enzymol.* 303:19-44 (1999)

Please visit our web site (<http://genome.rtc.riken.go.jp>) for further details.

FEATURES	SOURCE
Location/Qualifiers	1..302
/organism="Mus musculus"	
/strain="C57BL/6J"	
/db_xref="taxon:10090"	
/clone="4833401E05"	
/clone_1lb="RIKEN full-length enriched, 0 day neonate head"	
/sex="mixed"	
/tissue_type="head"	
/dev_stage="0 day neonate"	
/lab_host="DH10B"	
/note="Site_1: SalI; Site_2: BamHI. cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5', GAGAGAGAGACATCCCAAGCTCTTTTGTGTTTTTTTNN 3']. cDNA was prepared by using triethanolamine thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 10.0 and subtraction to Rot = 100.0. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAGAGATTTCGACTTAAATTAATCCCCCCCCCCC 3']. cDNA was cloned into the xhoI and BamHI sites. Vector: a modified pluscript KS(+) after bulk excision from Lambda FIC I."	
BASE COUNT	86 a 38 c 62 g 96 t
ORIGIN	
Query Match	39.6%; Score 35.6; DB 10; Length 302;
Best Local Similarity	64.6%; Pred. No. 8.9e+02;
Matches	53; Conservative 0; Mismatches 23; Indels 0; Gaps 0.
Oy	4 aaaaaattatccaagttctactgtaatttatgttcacgtttcatttgtaga 63 Db 185 AAAAAAATAATCATGACGATGAATTAATTGTTGCTGCCTCATCTTTGGCTGGGTTAGA 126 Oy 64 gcaataagtccaatgaagaaca 85 Db 125 CCTAGAGCTCGGGGATAGCAA 104
RESULT 3	
LOCUS	B67487 447 bp DNA GSS 09-DEC-1997
DEFINITION	T25M7R TAMU Arabidopsis thaliana genomic clone T25M7, DNA sequence.
ACCESSION	B67487
VERSION	B67487.1 GI:2666241
KEYWORDS	GSS
SOURCE	thale cress.
ORGANISM	Arabidopsis thaliana
	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis. 1 (bases 1 to 447)
REFERENCE	Rounsley,S.D., Field,C.E., Bass,S., Linher,K., Linher,K., Golden,K, Berry,K., Granger,D., Suh,E., Wible,C., Adams,M.D. and Venter ,J.C.
AUTHORS	A BAC End Sequence Database for Identifying Minimal Overlaps in Arabidopsis Genomic Sequencing. Update 3 Unpublished (1997) Contact: Steve Rounsley Department of Eukaryotic Genomics The Institute for Genomic Research
JOURNAL	
COMMENT	

9712 Medical Center Dr., Rockville, MD 20850, USA
Tel.: 301 838 0200
Fax: 301 838 0208
Email: roundsleyet@fcr.org
Seq primer: M13 Reverse
Class: BAC ends
High quality sequence stop: 447.

FEATURES

Source

```

/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="T25M7"
/clone_1b="TAMU"
/sex="hermaphrodite"
/notes="Vector: BelosacII; Site_1: HindIII; Site_2: HindIII
; Produced by Rod Wing"
BASE COUNT      187 a      64 c      64 g      132 t
ORIGIN

```

query Match	38.7%;	Score 34.8;	DB 13;	Length 447;
Best Local Similarity	65.4%;	Pred. No. 1e+03;		
Matches 51; Conservative	0;	Mismatches 27;	Indels 0;	Gaps 0;

OY 5 aaaaattatctcaagttcactgtaatttctagtcttcacgtttctctatgttgag 64
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 379 AAATAAAAATTGAAGTTTGATTCTTCGCAATTAATTTTTCTTCGTGGGG 320

QY	65	caataagtc	caatgaa	82
Db	319	TTCTGAAAAA	ACGGAAG	302

RESULT	4
LOCUS	AO968516/c
DEFINITION	AO968516 482 bp DNA
	ERJ037 ERG Arabidopsis thaliana genomic clone ERJ03, DNN
	28-JAN-2000

sequence.	
ACCESSION	AQ968516
VERSION	AQ968516.1
KEYWORDS	GI:6796217
	GSS.

ORGANISM

REFERENCE
1 (bases 1 to 482)

TITLE

JOURNAL
COMMENT

The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel.: 301 838 0200
Fax: 301 838 0208
Email: atc@tigr.org
For additional information, see <http://www.tigr.org/tdb/at/at.html>
Similar to A. thaliana chloroplast sequence (GB:AF000423)
Seq primer: TF
Class: shotgun.

FEATURES

Source

Location/Qualifiers
1..482
/organism="Arabidopsis thaliana"
/strain="Landsberg erecta"
/db_xref="taxon:3702"
/clone="LERJ03"
/clone_1bp="ERG"
/note="Organ: Leaf, Vector: pUC19JK, Total genomic DNA was sheared to 0.4-0.7 kbp before ligation."

BASE COUNT	208 a	64 c	78 g	132 t
ORIGIN				
Query Match	38.7%	Score 34.8:	DB 13:	Length 482:
Best Local Similarity	65.4%:	Pred. No. 1e+03:	27:	Indels 0:
Matches 51:	Conservative	0:	Mismatches	Gaps 0:

Oy 5 aaaaattatctcaagtttactgtaaattttatgcttcacatgcttcttatgttgag 64
||| ||| ||| ||| | ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 217 AAAATAAAATTTGGAAGTTTGATTCTTCCTTCAATTAATTTTTCTCTGTGGGG 158

QY	65	caataagtc	caatgaa	82
Db	157	TTCTGAAAAA	AAGAAG	140

RESULT	5			
LOCUS	A0968397/c			
DEFINITION	A0968397	488 bp	DNA	28-JAN-2000
	LEUC21TR	LERG	Arabidopsis thaliana	
			genomic clone	
			LEUC21, DNA	

ACCESSION	AQ968397
VERSION	AQ968397.1
KEYWORDS	GSS.

ORGANISM

REFERENCE
1 (bases 1 to 488)

TITLE

JOURNAL
COMMENT

The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: atef@tigr.org
For additional information, see <http://www.tigr.org/tdb/at/at.html>
Similar to A. thaliana chloroplast sequence (GB:AF000423)
Seq primer: TR
Class: shotgun.

FEATURES

Source

BASE COUNT	ORIGIN
186 a	78 c 83 g 141 t

Query Match	38.7%;	Score 34.8;	DB 13;	Length 488;
Best Local Similarity	65.4%;	Pred. No. 1e+03;		
Matches	51;	Conservative	0;	Mismatches 27;
			Indels	0;
			Gaps	0;

Qy 5 aaaaaattattccaagttactgtaatttatagtlcttcacagtctcatttgtagag 64
||| ||| ||| ||| | ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 95 AAATAAAATTTGAACAGTTTTGATTCTTCTTCGTAATTAATTTTTCTTCTGTGGGG 36

QY	65	caataagtc	caatgaag	82
Db	35	TTCTGAAAAA	AGAGAAG	18

RESULT 6
 LOCUS AO958266 497 bp DNA GSS 28-JAN-2000
 DEFINITION LERAW4/TR LERA Arabidopsis thaliana genomic clone LERAW47, DNA sequence.
 ACCESSION AO958266
 VERSION AO958266.1 GI:6785967
 KEYWORDS GSS.
 SOURCE thale cress.
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 REFERENCE 1 (bases 1 to 497)
 Buell, C.R., Lin, X., Pal, G., Barnstead, M., Bowman, C., Utterbach, T., Feldblum, T., Liang, F., Creasy, T., and Fraser, C.M.
 Genomic survey sequencing of Landsberg erecta ecotype of Arabidopsis thaliana and identification of sequence-based polymorphisms
 Unpublished (2000)
 JOURNAL COMMENT Contact: Xiaoying Lin
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: atetigr.org
 For additional information, see <http://www.tigr.org/tdb/at/at.html>
 Similar to A. thaliana chloroplast sequence (GB:AF00423)
 Seq primer: TR
 Class: shotgun.
 FEATURES
 source Location/Qualifiers
 1..497
 /organism="Arabidopsis thaliana"
 /strain="Landsberg erecta"
 /db_xref="taxon:3702"
 /clone="LERAW47"
 /clone_lib="LERA"
 /note="Organ: Leaf; Vector: PHOS1; Total genomic DNA was sheared to 0.9-1 kbp before ligation."
 BASE COUNT 144 a 82 c 69 g 202 t
 ORIGIN

Query Match 38.7%; Score 34.8; DB 13; Length 497;
 Best Local Similarity 65.4%; Pred. No. 9.9e+02;
 Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

OY 5 aaaaattatttcaagttaatttatttatttcaatgtttcattgtttgag 64
 ||| || |||| ||||| || |||| || |||| ||||| ||||| ||
 DB 337 AATATTAATTTGAAAGTTTCTTCCTGATTAATTTTCTTCTGCGG 396
 ||| || |||| ||||| || |||| || |||| ||||| ||||| ||
 OY 65 caataagtcgaatgaag 82
 | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 DB 397 TTCTGAAAAAAGAAAG 414

RESULT 7
 LOCUS B77731 498 bp DNA GSS 16-JAN-1998
 DEFINITION T28N11/TR TAMU Arabidopsis thaliana genomic clone T28N11, DNA sequence.
 ACCESSION B77731
 VERSION B77731.1 GI:2774370
 KEYWORDS GSS.
 SOURCE thale cress.
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 REFERENCE 1 (bases 1 to 498)
 Rounsley, S.D., Field, C.E., Bass, S., Linher, K., Linher, K., Golden, K., Berry, K., Granger, D., Suh, E., Wible, C., Adams, M.D. and Venter, J.C.

TITLE A BAC End Sequence Database for Identifying Minimal Overlaps in Arabidopsis Genomic Sequencing. Update 3
 JOURNAL UNPUBLISHED (1997)
 COMMENT Other_GSSs: T28N11/TR
 Contact: Steve Rounsley
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: rounsley@tigr.org
 Seq primer: M13-21
 Class: BAC ends
 High quality sequence stop: 498.
 FEATURES
 source Location/Qualifiers
 1..498
 /organism="Arabidopsis thaliana"
 /strain="Columbia"
 /db_xref="taxon:3702"
 /clone="T28N11"
 /clone_lib="TAMU"
 /sex="hermaphrodite"
 /note="Vector: BclBACII; Site_1: HindIII; Site_2: HindIII; Produced by Rod Wing"
 BASE COUNT 210 a 68 c 72 g 148 t
 ORIGIN

Query Match 38.7%; Score 34.8; DB 13; Length 498;
 Best Local Similarity 65.4%; Pred. No. 9.9e+02;
 Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

OY 5 aaaaattatttcaagttaatttatttatttcaatgtttcattgtttgag 64
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 DB 379 AATATTAATTTGAAAGTTTCTTCCTGATTAATTTTCTTCTGCGG 320
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 OY 65 caataagtcgaatgaag 82
 | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 DB 319 TTCTGAAAAAAGAAAG 302

RESULT 8
 LOCUS B30114 623 bp DNA GSS 13-OCT-1997
 DEFINITION T26F22/TRB TAMU Arabidopsis thaliana genomic clone T26F22, DNA sequence.
 ACCESSION B30114
 VERSION B30114.1 GI:2516080
 KEYWORDS GSS.
 SOURCE thale cress.
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 REFERENCE 1 (bases 1 to 623)
 Rounsley, S.D., Kelley, J.M., Field, C.E., Craven, M.B., Adams, M.D. and Venter, J.C.
 Use of a BAC End Sequence Database To Identify Minimal Overlaps for Arabidopsis Genomic Sequencing.
 Unpublished (1997)
 Other_GSSs: T26F22/TRB
 Contact: Steve Rounsley
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: rounsley@tigr.org
 Seq primer: M13-21
 Class: BAC ends
 High quality sequence stop: 623.
 FEATURES
 source Location/Qualifiers
 1..623

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/organism="Arabidopsis thaliana"
/strain="Columbia"
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/clone_1id="TAMU"
/sex="hermaphrodite"
/note="Vector: BelobACII; Site_1: HindIII; Site_2: HindIII
; Produced by Rod Wing"
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ORIGIN

Query Match      38.7%; Score 34.8; DB 13; Length 623;
Best Local Similarity 65.4%; Pred. No. 8.9e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

Oy 5 aaaaattattccaagttctgaatttctatgtttcttcttcttcttctgag 64
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 378 AAATTAATAATTGAAGATTCTCTCTCGAATTATTTCTCTCTGTGGG 319
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Oy 65 caataagtcgaatgaag 82
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 318 TTCTGAAAAAAGAAAG 301
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 9
B78178/c 627 bp DNA GSS 16-JAN-1998
LOCUS T3J11TF TAMU Arabidopsis thaliana genomic clone T3J11, DNA
DEFINITION
sequence.
ACCESSION B78178
VERSION B78178.1 GI:2774817
KEYWORDS GSS.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 627)
Rounsley,S.D., Field,C.E., Bass,S., Linher,K., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., Adams,M.D. and Venter
,J.C.
A BAC End Sequence Database for Identifying Minimal Overlaps in
Arabidopsis Genomic Sequencing. Update 3
JOURNAL Unpublished (1997)
COMMENT Other-GSS: T3J11TR
Contact: Steve Rounsley
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: rounsley@tigr.org
Seq primer: M13-21
Class: BAC ends
High quality sequence stop: 627.
Location/Qualifiers
1. 627
/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
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/sex="hermaphrodite"
/note="Vector: BelobACII; Site_1: HindIII; Site_2: HindIII
; Produced by Rod Wing"
BASE COUNT      261 a      87 c      100 g      178 t      1 others
ORIGIN

Query Match      38.7%; Score 34.8; DB 13; Length 627;
Best Local Similarity 65.4%; Pred. No. 8.9e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

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Oy 5 aaaaattattccaagttctgaatttctatgtttcttcttcttcttctgag 64
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Db 371 AAATTAATAATTGAAGATTCTCTCTCGAATTATTTCTCTCTGTGGG 312
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Oy 65 caataagtcgaatgaag 82
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 311 TTCTGAAAAAAGAAAG 294
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RESULT 10
A0958844 640 bp DNA GSS 28-JAN-2000
LOCUS LEREF16TR LERE Arabidopsis thaliana genomic clone LEREF16, DNA
DEFINITION
sequence.
ACCESSION A0958844
VERSION A0958844.1 GI:6786545
KEYWORDS GSS.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 640)
Buell,C.R., Lin,X., Pal,G., Barnstead,M., Bowman,C., Utterbach,T.,
Feldblyum,T., Jiang,F., Creasy,T. and Fraser,C.M.
Genomic survey sequencing of Landsberg erecta ecotype of
Arabidopsis thaliana and identification of sequence-based
polymorphisms
Unpublished (2000)
JOURNAL Contact: Xiaoying Lin
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: at@tigr.org
For additional information, see http://www.tigr.org/tdb/at/at.html
Similar to A. thaliana chloroplast sequence (GB:AP000423)
Seq primer: TR
Class: Shotgun.
FEATURES
source
1. 640
/organism="Arabidopsis thaliana"
/strain="Landsberg erecta"
/db_xref="taxon:3702"
/clone="LEREF16"
/clone_1id="LERE"
/note="Organ: Leaf; Vector: pUC19TK; Total genomic DNA was
sheared to 0.6-0.8 kbp before ligation."
BASE COUNT      181 a      99 c      83 g      277 t
ORIGIN

Query Match      38.7%; Score 34.8; DB 13; Length 640;
Best Local Similarity 65.4%; Pred. No. 8.8e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

Oy 5 aaaaattattccaagttctgaatttctatgtttcttcttcttcttctgag 64
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Db 147 AAATTAATAATTGAAGATTCTCTCTCGAATTATTTCTCTCTGTGGG 206
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Oy 65 caataagtcgaatgaag 82
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 207 TTCTGAAAAAAGAAAG 224
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RESULT 11
B98363/c 643 bp DNA GSS 31-MAR-1998
LOCUS T24C11TFB TAMU Arabidopsis thaliana genomic clone T24C11, DNA
DEFINITION
sequence.
ACCESSION B98363
VERSION B98363.1 GI:3000442

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[illegible]

```

BP 191 91006 EVRY cedex - FRANCE (E-mail: seqref@genome.cns.fr)
- Web : www.genome.cns.fr

FEATURES
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                /strain="Columbia"
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                /clone_lib="TAMU"
                /clone="T1202"
                /note="end : 77"

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ORIGIN

Query Match      38.7% Score 34.8: DB 13: Length 660;
Best Local Similarity 65.4%: Pred. No. 8.7e+02;
Matches 51: Conservative 0: Mismatches 27: Indels 0: Gaps 0:

OY      5 aaaaattatcgaagttacgtgaatttatagttcttcattgttgag 64
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      371 AATATAAATTGGAAGTTTGATTTCTCTTCATTAATTTTCTCTGTGGG 312

OY      65 caatagcgaatgaag 82
      | | | | |
Db      311 TTCTGAAAAAAGCAAG 294

RESULT 13
A0968517      683 bp      DNA      GSS      28-JAN-2000
LOCUS      LERJ03TR LERG Arabidopsis thaliana genomic clone LERJ03, DNA
DEFINITION      sequence.
ACCESSION      A0968517
VERSION      A0968517.1 GI:6796218
KEYWORDS      GSS.
SOURCE      thale cress.
ORGANISM      Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE      1 (bases 1 to 683)
AUTHORS      Buell,C.R., Lin,X., Pat,G., Barnstead,M., Bowman,C., Uterbach,T.,
      Feldblyum,T., Liang,F., Creasy,T. and Fraser,C.M.
      Genomic survey sequencing of Landsberg erecta ecotype of
      Arabidopsis thaliana and identification of sequence-based
      polymorphisms
      Unpublished (2000)
JOURNAL
COMMENT      Contact: Xiaoying Lin
      The Institute for Genomic Research
      9712 Medical Center Dr., Rockville, MD 20850, USA
      Tel: 301 838 0200
      Fax: 301 838 0208
      Email: atetlgr.org
      For additional information, see http://www.tigr.org/cdb/at.html
      Similar to A. thaliana chloroplast sequence (GB:AP000423)
      Seq primer: TR
      Class: Shotgun.
FEATURES
    source
        Location/Qualifiers
            1..683
                /organism="Arabidopsis thaliana"
                /strain="Landsberg erecta"
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                /clone="LERJ03"
                /clone_lib="LENG"
                /note="Organ: Leaf; Vector: pUC19JK; Total genomic DNA was
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BASE COUNT      208 a      111 c      88 g      276 t

ORIGIN

Query Match      38.7% Score 34.8: DB 13: Length 683;
Best Local Similarity 65.4%: Pred. No. 8.5e+02;
Matches 51: Conservative 0: Mismatches 27: Indels 0: Gaps 0:

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 QY 65 caataagtcacatgaag 82
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 Db 572 TTCTGAAAAAAGAAAG 589

RESULT 14
 AO958265 743 bp DNA GSS 28-JAN-2000
 LOCUS LERRA47TF LERRA Arabidopsis thaliana genomic clone LERRA47, DNA
 DEFINITION sequence.
 ACCESSION AO958265
 VERSION AO958265.1 GI:6785966
 KEYWORDS GSS.
 SOURCE thale cress.
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 1 (bases 1 to 743)
 Buell,C.R., Lin,X., Pal,G., Barnstead,M., Bowman,C., Uterbach,T.,
 Feldblyum,T., Jiang,F., Creasy,T. and Fraser,C.M.
 Genomic survey sequencing of Landsberg erecta ecotype of
 Arabidopsis thaliana and identification of sequence-based
 polymorphisms
 JOURNAL Unpublished (2000)
 COMMENT Contact: Xiaoying Lin
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: atcligr.org
 For additional information, see <http://www.tigr.org/cdb/at.html>
 Similar to A. thaliana chloroplast sequence (GB:AP000423)
 Seq primer: TF
 Class: Shotgun.

FEATURES
 source
 Location/Qualifiers
 1..743
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 Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

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 ||| || |||| ||||| | ||| || ||||| ||||| |
 Db 534 AAATTAATAATTGAAAGTTTGATTTCTTCTTGCAATTAATTTTCTTGTGGG 475

QY 65 caataagtcacatgaag 82
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 Db 474 TTCTGAAAAAAGAAAG 457

RESULT 15
 AO956484 798 bp DNA GSS 28-JAN-2000
 LOCUS LERRA26TR LERRA Arabidopsis thaliana genomic clone LERRA26, DNA
 DEFINITION sequence.
 ACCESSION AO956484
 VERSION AO956484.1 GI:6784185

KEYWORDS GSS.
 SOURCE thale cress.
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 1 (bases 1 to 798)
 Buell,C.R., Lin,X., Pal,G., Barnstead,M., Bowman,C., Uterbach,T.,
 Feldblyum,T., Jiang,F., Creasy,T. and Fraser,C.M.
 Genomic survey sequencing of Landsberg erecta ecotype of
 Arabidopsis thaliana and identification of sequence-based
 polymorphisms
 JOURNAL Unpublished (2000)
 COMMENT Contact: Xiaoying Lin
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: atcligr.org
 For additional information, see <http://www.tigr.org/cdb/at.html>
 Similar to A. thaliana chloroplast sequence (GB:AP000423)
 Seq primer: TR
 Class: Shotgun.

FEATURES
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 Location/Qualifiers
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 BASE COUNT 231 a 115 c 109 g 343 t
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Query Match 38.7%; Score 34.8; DB 13; Length 798;
 Best Local Similarity 65.4%; Pred. No. 7.9e+02;
 Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

QY 5 aaaaattattccaagttctacgttaattttatgtttcatttcttattgttgag 64
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 Db 164 AAATTAATAATTGAAAGTTTGATTTCTTCTTGCAATTAATTTTCTTGTGGG 223

QY 65 caataagtcacatgaag 82
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 Db 224 TTCTGAAAAAAGAAAG 241

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 Job time: 2485 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:18:46 ; Search time 90.4 Seconds

(Without alignments)
225.476 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
Sequence: 1 atgaaaaaattatttcaaa.....gtccaatgaaagcaagtgc 90

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 351203 seqs, 113238999 residues

Total number of hits satisfying chosen parameters: 702406

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

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5: /cgn2_6/ptodata/2/ina/PCBUS.COMB.seq:.*
6: /cgn2_6/ptodata/2/ina/backfiles1.seq:.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	31.2	34.7	3337	1 US-08-072-610-1	Sequence 1, Appl
C 2	31.2	34.7	3337	2 US-08-719-822B-1	Sequence 1, Appl
C 3	31.2	34.7	3337	4 US-09-092-458-1	Sequence 1, Appl
C 4	30.6	34.0	87350	3 US-08-781-891-79	Sequence 79, Appl
C 5	30.0	33.3	4140	3 US-08-884-731-2	Sequence 2, Appl
C 6	29.6	32.9	1703	4 US-08-378-313-18	Sequence 18, Appl
C 7	28.2	31.3	2334	1 US-08-062-632-4	Sequence 4, Appl
C 8	28	31.1	1013	1 US-07-920-519-30	Sequence 30, Appl
C 9	28	31.1	1013	1 US-08-086-410-23	Sequence 23, Appl
C 10	28	31.1	1013	1 US-08-314-586-30	Sequence 30, Appl
C 11	27.8	30.9	1584	1 US-07-667-376A-1	Sequence 1, Appl
C 12	27.4	30.4	1100	4 US-07-861-458C-4	Sequence 4, Appl
C 13	27.4	30.4	5613	2 US-08-463-418-1	Sequence 1, Appl
C 14	27.2	30.2	602	1 US-08-764-100-8	Sequence 8, Appl
C 15	27.2	30.2	642	1 US-08-764-100-13	Sequence 13, Appl
C 16	27.2	30.2	662	1 US-08-764-100-7	Sequence 7, Appl
C 17	27.2	30.2	662	1 US-08-998-416-185	Sequence 185, App
C 18	27.2	30.2	663	4 US-08-998-416-191	Sequence 191, App
C 19	27.2	30.2	663	4 US-08-998-416-937	Sequence 937, App
C 20	27.2	30.2	701	4 US-08-998-416-701	Sequence 701, App
C 21	27.2	30.2	711	4 US-08-998-416-786	Sequence 786, App
C 22	27.2	30.2	724	4 US-08-998-416-683	Sequence 683, App
C 23	27.2	30.2	732	4 US-08-998-416-1036	Sequence 1036, App
C 24	27.2	30.2	767	4 US-08-998-416-472	Sequence 472, App
C 25	27.2	30.2	782	4 US-08-998-416-224	Sequence 224, App
C 26	27.2	30.2	827	4 US-08-998-416-535	Sequence 535, App
C 27	27.2	30.2	828	4 US-08-998-416-538	Sequence 538, App

C 28	27.2	30.2	834	4 US-08-998-416-305	Sequence 305, App
C 29	27.2	30.2	854	4 US-08-998-416-534	Sequence 534, App
C 30	27.2	30.2	860	4 US-08-998-416-287	Sequence 287, App
C 31	27.2	30.2	2993	1 US-08-764-100-2	Sequence 2, Appl
C 32	27.2	30.2	2993	1 US-08-764-100-10	Sequence 10, Appl
C 33	27.2	30.2	3000	1 US-08-764-100-9	Sequence 9, Appl
C 34	27.2	30.2	3001	1 US-08-764-100-1	Sequence 1, Appl
C 35	27.2	30.2	14176	1 US-08-307-499-1	Sequence 1, Appl
C 36	27.2	30.2	14176	1 US-08-307-499-14	Sequence 14, Appl
C 37	27.2	30.2	14176	4 US-09-299-268-1	Sequence 1, Appl
C 38	27.2	30.2	14176	4 US-09-299-268-14	Sequence 14, Appl
C 39	27	30.0	642	3 US-08-817-926-49	Sequence 49, Appl
C 40	27	30.0	839	3 US-08-817-926-50	Sequence 50, Appl
C 41	27	30.0	1478	3 US-08-817-926-1	Sequence 1, Appl
C 42	27	30.0	1478	4 US-09-297-053-2	Sequence 2, Appl
C 43	27	30.0	3562	3 US-08-817-926-19	Sequence 19, Appl
C 44	26.8	29.8	2168	3 US-08-749-522-6	Sequence 6, Appl
C 45	26.8	29.8	7083	4 US-09-198-839-1	Sequence 1, Appl

ALIGNMENTS

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RESULT 1
US-08-072-610-1/c
: Sequence 1, Application US/08072610
: Patent No. 5532133
:
: GENERAL INFORMATION:
: APPLICANT: Barnwell, John
: TITLE OF INVENTION: Plasmidium vivax Blood Stage Antigens,
: TITLE OF INVENTION: Monoclonal Antibodies, and Diagnostic Assays
: NUMBER OF SEQUENCES: 2
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Darby and Darby
: STREET: 805 Third Ave.
: CITY: New York
: STATE: New York
: COUNTRY: USA
: ZIP: 10022-7513
:
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.25
:
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/072,610
: FILING DATE: 19930602
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: Gogoris, Adga
: REGISTRATION NUMBER: 29,714
: REFERENCE/DOCKET NUMBER: 5986/07686
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (212)527-7700
: TELEFAX: (212)753-6237
: TELEX: 236687
:
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 3337 base pairs
: TYPE: NUCLEIC ACID
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: HYPOTHETICAL: NO
: ANTI-SENSE: NO
: ORIGINAL SOURCE:
: ORGANISM: Plasmidium vivax
: IMMEDIATE SOURCE:
: CLONE: pVMB3.3.1
:
: US-08-072-610-1
:
: Query Match 34.7%; Score 31.2; DB 1; Length 3337;
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Best Local Similarity 63.2%; Pred. No. 5.3;
Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;

QY 10 attatttcaaaagttactcgttaattttatgtttcatttcttctatgttggaagcaata 69

Db 138 AATATATATGTAATATAGCAGCTTTTATAGATATTTTATCTTACTGTAACACATGA 79

QY 70 agtccaatgaagcaaa 85

Db 78 ATTAAGAAAGCAAA 63

RESULT 2

US-08-719-822B-1/c

; Sequence 1, Application US/08719822B

; Patent No. 5874527

; GENERAL INFORMATION:

; APPLICANT: Barnwell, John

; TITLE OF INVENTION: Plasmodium vivax Blood Stage Antigens

; NUMBER OF SEQUENCES: 4

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Darby and Darby

; STREET: 805 Third Ave.

; CITY: New York

; STATE: New York

; COUNTRY: USA

; ZIP: 10022-7513

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/719,822B

; FILING DATE: 09/30/96

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Gogoris, Agda

; REGISTRATION NUMBER: 29,714

; REFERENCE/DOCKET NUMBER: 5986/176860S2

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (212)527-7700

; TELEFAX: (212)753-6237

; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 3337 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: double

; TOPOLOGY: linear

; MOLECULE TYPE: DNA (genomic)

; HYPOTHETICAL: NO

; ANTI-SENSE: NO

; ORIGINAL SOURCE:

; ORGANISM: Plasmodium vivax

; IMMEDIATE SOURCE:

; CLONE: pVMB3.3.1

; US-08-719-822B-1

Query Match 34.7%; Score 31.2; DB 2; Length 3337;

Best Local Similarity 63.2%; Pred. No. 5.3;

Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;

QY 10 attatttcaaaagttactcgttaattttatgtttcatttcttctatgttggaagcaata 69

Db 138 AATATATATGTAATATAGCAGCTTTTATAGATATTTTATCTTACTGTAACACATGA 79

QY 70 agtccaatgaagcaaa 85

Db 78 ATTAAGAAAGCAAA 63

RESULT 3

US-09-092-458-1/c

; Sequence 1, Application US/09092458

; Patent No. 6231861

; GENERAL INFORMATION:

; APPLICANT: Barnwell, John

; TITLE OF INVENTION: Plasmodium vivax Blood Stage Antigens, Monoclonal Antibodies, and Diagnostic Assays

; NUMBER OF SEQUENCES: 4

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Darby and Darby

; STREET: 805 Third Ave.

; CITY: New York

; STATE: New York

; COUNTRY: USA

; ZIP: 10022-7513

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/092,458

; FILING DATE:

; CLASSIFICATION:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/719,821

; FILING DATE: 09/30/96

; ATTORNEY/AGENT INFORMATION:

; NAME: Gogoris, Agda

; REGISTRATION NUMBER: 29,714

; REFERENCE/DOCKET NUMBER: 5986/176860S3

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (212)527-7700

; TELEFAX: (212)753-6237

; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 3337 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: double

; TOPOLOGY: linear

; MOLECULE TYPE: DNA (genomic)

; HYPOTHETICAL: NO

; ANTI-SENSE: NO

; ORIGINAL SOURCE:

; ORGANISM: Plasmodium vivax

; IMMEDIATE SOURCE:

; CLONE: pVMB3.3.1

; US-09-092-458-1

Query Match 34.7%; Score 31.2; DB 4; Length 3337;

Best Local Similarity 63.2%; Pred. No. 5.3;

Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;

QY 10 attatttcaaaagttactcgttaattttatgtttcatttcttctatgttggaagcaata 69

Db 138 AATATATATGTAATATAGCAGCTTTTATAGATATTTTATCTTACTGTAACACATGA 79

QY 70 agtccaatgaagcaaa 85

Db 78 ATTAAGAAAGCAAA 63

RESULT 4

US-08-781-891-79

; Sequence 79, Application US/08781891

; Patent No. 6090620

; GENERAL INFORMATION:

; APPLICANT: Fu, Ying-Hui

; APPLICANT: Yu, Chang-En

; APPLICANT: Oshima, Junko


```

Query Match          34.0%; Score 30.6; DB 3; Length 87350;
Best Local Similarity 62.3%; Pred. NO. 9;
Matches 48; Conservative 0; Mismatches 29; Indels 0; Gaps 0;

QY      1 atgaaaaaatattctcaagttctactgcaattttatctgattcttcattglt 60
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 76835 ATGCAATAGGGGTATTTGCAAGSTTCTGTATGTATTCTGTAGAAAGTATCTCAAGGG 76894

QY      61 ggagcaataagttccaat 77
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 76895 GGATATATATCTTCATT 76911

RESULT      5
US-08-894-731-2
: Sequence 2, Application US/08894731
: Patent No. 6084089
: GENERAL INFORMATION:
: APPLICANT: MINE, Toshiki
: APPLICANT: OHYAMA, Akio
: APPLICANT: HIYOSHI, Toru
: APPLICANT: KASAKOKA, Keisuke
: TITLE OF INVENTION: COLD-INDUCIBLE PROMOTER SEQUENCE
: FILE REFERENCE: 760-234P
: CURRENT APPLICATION NUMBER: US/08/894,731
: CURRENT FILING DATE: 1997-10-27
: NUMBER OF SEQ ID NOS: 8
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 2
: LENGTH: 4140
: TYPE: DNA
: ORGANISM: Solanum tuberosum
US-08-894-731-2

Query Match          33.3%; Score 30; DB 3; Length 4140;

```

```

US-08-378-313-18      RESULT        6
US-08-378-313-18      : Sequence 18, Application US/08378313
                        : Patent No. 6207881
                        :
                        : GENERAL INFORMATION:
                        : APPLICANT: THEOLOGIS, ATHANASIOS
                        : APPLICANT: SATO, TAKAHIDO
                        : TITLE OF INVENTION: CONTROL OF FRUIT RIPENING THROUGH
                        : TITLE OF INVENTION: GENETIC CONTROL OF ACC SYNTHASE SYNTHESIS
                        : NUMBER OF SEQUENCES: 34
                        : CORRESPONDENCE ADDRESS:
                        : ADDRESSEE: MORRISON & FOERSTER
                        : STREET: 755 Page Mill Road
                        : CITY: Palo Alto
                        : STATE: California
                        : COUNTRY: USA
                        : ZIP: 94304-1018
                        :
                        : COMPUTER READABLE FORM:
                        : MEDIUM TYPE: Floppy disk
                        : COMPUTER: IBM PC compatible
                        : OPERATING SYSTEM: PC-DOS/MS-DOS
                        : SOFTWARE: Patentin Release #1.0, Version #1.25
                        : CURRENT APPLICATION DATA:
                        : APPLICATION NUMBER: US/08/378,313
                        : FILING DATE:
                        : CLASSIFICATION: 800
                        : PRIOR APPLICATION DATA:
                        : APPLICATION NUMBER: US 07/862,493
                        : FILING DATE: 02-APR-1992
                        : ATTORNEY/AGENT INFORMATION:
                        : NAME: MURASHIGE, KATE H.
                        : REGISTRATION NUMBER: 29,959
                        : REFERENCE/DOCKET NUMBER: 29190-20002.20
                        : TELECOMMUNICATION INFORMATION:
                        : TELEPHONE: (415) 856-5600
                        : TELEFAX: (415) 494-0792
                        : TELEX: 706141
                        : INFORMATION FOR SEQ ID NO: 18:
                        : SEQUENCE CHARACTERISTICS:
                        : LENGTH: 1703 base pairs
                        : TYPE: nucleic acid
                        : STRANDEDNESS: single
                        : TOPOLOGY: linear
                        : FEATURE:
                        : NAME/KEY: CDS
                        : LOCATION: 11..1489
                        :
US-08-378-313-18

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Query Match	32.9%	Score 29.6;	DB 4;	Length 1703;
Best Local Similarity	64.7%;	Pred. No. 12;		
Matches 44;	Conservative 0;	Mismatches 24;	Indels 0;	Gaps 0;
OY	1 atgaaaaaatatttccaagtctcgttaattttatgttttcacgtttctattgtt 60			
Db	1492 atttaaaacattttccaamaatttcataccattcacatattgttttttttttt 1551			
OY	61 ggaagcaat 68			
Db	1552 gggtcatt 1559			

RESULT 7
US-08-062-632-4/C
; Sequence 4, Application US/08062632
; Patent No. 5712090
; GENERAL INFORMATION:
; APPLICANT: Artushin, Sergey
; APPLICANT: Stipkovits, Laslo
; APPLICANT: Minion, F. Chris
; TITLE OF INVENTION: PCR-Based Assay For Mycoplasma
; TITLE OF INVENTION: Hypopneumoniae
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Dickstein, Shapiro and Morin
; STREET: 2101 L. St. NW
; CITY: Washington
; STATE: D.C.
; COUNTRY: USA
; ZIP: 22037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/062,632
; FILING DATE: 18-MAY-1993
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Brady Jr., James W.
; REGISTRATION NUMBER: 32,115
; REFERENCE/DOCKET NUMBER: 18900.018
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202)785-9700
; TELEFAX: (202)887-0689
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2334 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-062-632-4

Query Match 31.3%; Score 28.2; DB 1; Length 2334;
Best Local Similarity 64.6%; Pred. No. 28;
Matches 42; Conservative 0; Mismatches 23; Indels 0; Gaps 0;
QY 1 atgaaataattctcaagttactgtaattttatgtttcattcttattgtt 60
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 236 ATTAGAAATAACGGCTTATTGTGTATTATTATTATTACTTGCTTTAATAGTT 177
QY 61 ggaagc 65
|||
DB 176 TCTGC 172

RESULT 8
US-07-920-519-30/C
; Sequence 30, Application US/07920519
; Patent No. 5382518
; GENERAL INFORMATION:
; APPLICANT: CAPUT, DANIEL
; APPLICANT: FERRARA, PASQUAL
; APPLICANT: GUILLEMOT, JEAN-CLAUDE
; APPLICANT: KAGHAD, MOURAD
; APPLICANT: LEGOUX, RICHARD
; APPLICANT: LOISON, GERARD
; APPLICANT: LARRE, ELIZABETH
; APPLICANT: LUPKER, JOHANNES
; APPLICANT: LEPLATOIS, PASCUAL
; APPLICANT: SALOME, MARK

;; TITLE OF INVENTION: URATE OXIDASE ACTIVITY PROTEIN,
;; TITLE OF INVENTION: MICRO-ORGANISMS AND TRANSFORMED CELLS
;; TITLE OF INVENTION: MICRO-ORGANISMS AND TRANSFORMED CELLS
;; NUMBER OF SEQUENCES: 36
;; CORRESPONDENCE ADDRESSES:
;; ADDRESSEE: Foley & Lardner
;; STREET: 1800 Diagonal Road, Suite 500
;; CITY: Alexandria
;; STATE: Virginia
;; COUNTRY: USA
;; ZIP: 22313-0299
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: Patentin Release #1.0, Version #1.25
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/07/920,519
;; FILING DATE:
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US/07/659,408
;; FILING DATE:
;; ATTORNEY/AGENT INFORMATION:
;; NAME: BENT, Stephen A.
;; REGISTRATION NUMBER: 29,768
;; REFERENCE/DOCKET NUMBER: 16781/276 BEDL
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (703)836-9300
;; TELEFAX: (703)683-4109
;; TELEX: 899149
;; INFORMATION FOR SEQ ID NO: 30:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 1013 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; HYPOTHETICAL: NO
;; IMMEDIATE SOURCE:
;; CLONE: Fragment D
; US-07-920-519-30

Query Match 31.1%; Score 28; DB 1; Length 1013;
Best Local Similarity 60.5%; Pred. No. 30;
Matches 46; Conservative 0; Mismatches 30; Indels 0; Gaps 0;
QY 11 ttattcaagtttactgtaattttatgtttcattcttattgttggaacaataa 70
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 1004 TTTTNTTTTNTTTTNTTTTNTTTTNTTTTNTTTTNTTTTNTTTTNTTGAAGTAA 945
QY 71 gtcaatgaagaag 86
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 944 AACAGTGATGATGATG 929

RESULT 9
US-08-086-410-23/C
; Sequence 23, Application US/08086410
; Patent No. 5407822
; GENERAL INFORMATION:
; APPLICANT: LEPLATOIS, Pascal
; APPLICANT: LOISON, Gerard
; APPLICANT: PESSEQUE, Bernard
; APPLICANT: SHIRE, David
; TITLE OF INVENTION: Artificial promoter for the expression
; TITLE OF INVENTION: of proteins in yeast
; NUMBER OF SEQUENCES: 37
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: FOLEY & LARDNER
; STREET: King Street Station, Suite 500, 1800 Diagonal
; STREET: Road, PO Box 299

STREET: 3000 K Street, Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/314,586
FILING DATE: 28-SEP-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/659,408
FILING DATE: 25-APR-1991
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16781/509/BDL
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)836-9300
TELEFAX: (703)683-4109
TELEX: 899149
INFORMATION FOR SEQ ID NO: 30:
SEQUENCE CHARACTERISTICS:
LENGTH: 1013 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
IMMEDIATE SOURCE:
CLONE: Fragment D
US-08-314-586-30

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Query Match          31.1%; Score 28; DB 1; Length 1013;
Best Local Similarity 60.5%; Pred. NO. 30;
Matches 46; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

OY      11 ttattcaaaagttcactgttaattctatgcttcacgttctctatttgtagccaataa 70
       ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      1004 TTTT'TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTGGAAGTAATAA 945

OY      71 gtccaatgaagaacaag 86
       | | | | | | | |
Db      944 AACACGTGATGATGAATG 929

RESULT 11
US-07-667-276A-1
Sequence 1, Application US/07667276A
Patent No. 5470971
GENERAL INFORMATION:
APPLICANT: Kondo, Keiji
APPLICANT: Inouye, Masayori
TITLE OF INVENTION: STRESS-INDUCED PROTEINS, GENES CODING
TITLE OF INVENTION: THEREFOR, TRANSFORMED CELLS OF ORGANISMS, METHODS AND
TITLE OF INVENTION: APPLICATIONS
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Weisner & Associates
STREET: 230 S. Fifteenth Street, Suite 500
CITY: Philadelphia
STATE: PA
COUNTRY: USA
ZIP: 19102
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

```

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SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/667,276A
FILING DATE: 11-MAR-1991
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Weiser, Gerard J.
REGISTRATION NUMBER: 19,763
REFERENCE/DOCKET NUMBER: 377.5351P
TELECOMMUNICATION INFORMATION:
TELEPHONE: 215-875-8383
TELEFAX: 215-875-8394
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1584 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
FRAGMENT TYPE: Internal
ORIGINAL SOURCE:
ORGANISM: Saccharomyces cerevisiae
STRAIN: S288C
FEATURE:
NAME/KEY: CDS
LOCATION: 475..1104
FEATURE:
NAME/KEY: misc-feature
LOCATION: 1
OTHER INFORMATION: /note= "Base #1 of Sequence No. 5470971 1
OTHER INFORMATION: corresponds to base -474 of the sequence listed in
OTHER INFORMATION: Figure 4 of the application"
US-07-667-276A-1

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Query Match      30.9%; Score 27.8; DB 1; Length 1584;
Best Local Similarity 62.0%; Pred. No. 34;
Matches 44; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

```

```

QY 2 tgaataaatttcaagttactgtatatttattgttcaatgtttcttattgttg 61
    | | | | | | | | | | | | | | | | | | | | | | | | | | | |
DB 1119 TTACACCAATATTTGAAATATACCTCATATATATACCTTCTTATGTAATGTGA 1178
    | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 62 gagcaataagt 72
    | | | | | | |
DB 1179 GTTCGAAATTT 1189
    | | | | | | |

```

```

RESULT 12
US-07-861-458C-4/C
Sequence 4, Application US/07661458C
Patent No. 6232061
GENERAL INFORMATION:
APPLICANT: Marchionni, Mark Andrew
APPLICANT: Johnson, Carl D.
TITLE OF INVENTION: HOMOLOGY CLONING
NUMBER OF SEQUENCES: 142
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson
STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 MB
COMPUTER: IBM PS/2 Model 502 or 55SX
OPERATING SYSTEM: MS-DOS (Version 5.0)
SOFTWARE: WordPerfect (Version 5.1)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/861,458C

```

```

FILING DATE: 04/01/92
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Clark, Paul T.
REGISTRATION NUMBER: 30,162
REFERENCE/DOCKET NUMBER: 04585/014001
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1100
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-07-861-458C-4

```

```

Query Match      30.4%; Score 27.4; DB 4; Length 1100;
Best Local Similarity 59.7%; Pred. No. 42;
Matches 46; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

```

```

QY 11 ttattcaagttactgtaattttatgttttcaatgtttcttattgttgagcaataa 70
    | | | | | | | | | | | | | | | | | | | | | | | | | | | |
DB 1079 TTTTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 1020
    | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 71 gtccaatgaagcaagt 87
    | | | | | | |
DB 1019 GCTCAGTATATACATTT 1003
    | | | | | | |

```

```

RESULT 13
US-08-463-418-1
Sequence 1, Application US/08463418
Patent No. 5908971
GENERAL INFORMATION:
APPLICANT: Van Der Straeten, Dominique et al.
TITLE OF INVENTION: CROCIER ACC SYNTHASE AND USES THEREOF
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 225 Franklin Street
CITY: Boston
STATE: MA
COUNTRY: USA
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/463,418
FILING DATE: 05-JUN-1995
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/962,481
FILING DATE: 15-OCT-1992
ATTORNEY/AGENT INFORMATION:
NAME: Clark, Paul T.
REGISTRATION NUMBER: 30,162
REFERENCE/DOCKET NUMBER: 00786/161002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617/542-5070
TELEFAX: 617/542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5613 base pairs

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 03:24:24 ; Search time 93.51 Seconds
(without alignments)
217.977 Million cell updates/sec

Title: US-09-531-438-4

Sequence: 1 atgaaaaaatatttcaaa.....gtccaatgaagaagcagtga 90

Scoring table: OLIGO-NUC

Gapop 60.0, Gapext 60.0

Searched: 351203 seqs, 113238999 residues

Word size : 0

Total number of hits satisfying chosen parameters: 495388

Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database : Issued Patents - NA:*

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3: /cgn2_6/ptodata/2/ina/6A.COMB.seq:*
4: /cgn2_6/ptodata/2/ina/6B.COMB.seq:*
5: /cgn2_6/ptodata/2/ina/6C.COMB.seq:*
6: /cgn2_6/ptodata/2/ina/backfile1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	No.	Score	Query	Match	Length	DB	ID	Description
	1	13	14.4	24	3	US-08-672-215-1		Sequence 1, Appl
	2	13	14.4	30	2	US-08-629-001A-79		Sequence 79, Appl
	3	13	14.4	30	4	US-08-642-274D-158		Sequence 158, App
	4	13	14.4	38	5	PCT-US96-00547-40		Sequence 40, Appl
	5	13	14.4	39	1	US-08-105-483-168		Sequence 168, App
	6	13	14.4	39	1	US-08-709-209-168		Sequence 168, App
	7	13	14.4	39	1	US-08-303-275-56		Sequence 56, Appl
	8	13	14.4	39	1	US-08-458-101-168		Sequence 168, App
	9	13	13.3	14	1	US-08-271-880A-201		Sequence 201, App
	10	13	13.3	14	1	US-08-910-408-201		Sequence 201, App
	11	13	13.3	14	3	US-09-249-215-201		Sequence 201, App
	12	13	13.3	18	3	US-09-163-162-19		Sequence 19, Appl
	13	13	13.3	18	4	US-09-286-407-19		Sequence 19, Appl
	14	13	13.3	21	1	US-08-661-507-6		Sequence 6, Appl
	15	13	13.3	21	2	US-08-855-085-4		Sequence 4, Appl
	16	13	13.3	21	2	US-09-166-030-4		Sequence 4, Appl
	17	13	13.3	21	2	US-08-865-675-4		Sequence 4, Appl
	18	13	13.3	21	2	US-08-933-749-5		Sequence 5, Appl
	19	13	13.3	21	2	US-09-237-510-4		Sequence 4, Appl
	20	13	13.3	21	3	US-09-120-916-4		Sequence 4, Appl
	21	13	13.3	21	3	US-08-964-020-8		Sequence 8, Appl
	22	13	13.3	21	3	US-09-235-583-5		Sequence 5, Appl
	23	13	13.3	21	4	US-09-599-164-5		Sequence 5, Appl
	24	13	13.3	23	1	US-08-727-003A-10		Sequence 10, Appl
	25	13	13.3	23	3	US-08-487-799-11		Sequence 11, Appl
	26	12	13.3	24	2	US-08-210-762E-31		Sequence 31, Appl
	27	12	13.3	24	4	US-08-256-799-26		Sequence 26, Appl

28	12	13.3	24	4	US-08-462-437-26	Sequence 26, Appl
29	12	13.3	25	1	US-08-271-880A-140	Sequence 140, App
30	12	13.3	25	2	US-08-910-408-140	Sequence 140, App
31	12	13.3	25	3	US-09-249-215-140	Sequence 140, App
32	12	13.3	26	3	US-08-835-728D-72	Sequence 72, Appl
33	12	13.3	26	3	US-08-835-728D-72	Sequence 176, Appl
34	12	13.3	26	4	US-09-490-558-72	Sequence 176, Appl
35	12	13.3	26	4	US-09-490-558-176	Sequence 176, App
36	12	13.3	27	2	US-08-668-128B-6	Sequence 6, Appl
37	12	13.3	27	2	US-08-905-445-6	Sequence 6, Appl
38	12	13.3	27	3	US-08-644-116A-6	Sequence 6, Appl
39	12	13.3	28	2	US-08-771-602D-21	Sequence 21, Appl
40	12	13.3	28	2	US-08-771-602D-21	Sequence 20, Appl
41	12	13.3	30	1	US-08-619-724-3	Sequence 3, Appl
42	12	13.3	30	2	US-08-629-001A-25	Sequence 25, Appl
43	12	13.3	30	4	US-08-642-274D-104	Sequence 104, App
44	12	13.3	30	4	US-08-935-312-14	Sequence 14, Appl
45	12	13.3	33	1	US-08-341-456A-14	Sequence 14, Appl

ALIGNMENTS

RESULT 1
US-08-672-215-1/c
Sequence 1, Application US/08672215
Patent No. 6020121
GENERAL INFORMATION:
APPLICANT: Ying Bao, Amy Boggs, Pamela R. Contag,
APPLICANT: Nancy A. Federspiel, Alan Herbert,
APPLICANT: Scott J. Hecker, Francois Melouin
TITLE OF INVENTION: INHIBITORS OF REGULATORY PATHWAYS
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon 6 Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ for Windows 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/672.215
FILING DATE: June 25, 1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/004,626
FILING DATE: September 29, 1995
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 219/158
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 488-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-672-215-1

Query Match 14.4%; Score 13; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0;

QY 27 tgaattttatg 39
|||||
DB 20 tgaattttatg 8

RESULT 2

US-08-629-001A-79
; Sequence 79, Application US/08629001A
; Patent No. 5858661
; GENERAL INFORMATION:
; APPLICANT: Shiloh, Yosef
; TITLE OF INVENTION: ATAXIA-TELANGIECTASIA GENE AND ITS
; TITLE OF INVENTION: GENOMIC ORGANIZATION
; NUMBER OF SEQUENCES: 139
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kohn & Associates
; STREET: 30500 No. 5858661thwestern Hwy;
; CITY: Farmington Hills
; STATE: Michigan
; COUNTRY: US
; ZIP: 48334
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/629,001A
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Kohn, Kenneth I.
; REGISTRATION NUMBER: 30,955
; REFERENCE/DOCKET NUMBER: 2290,00032
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (810) 539-5050
; TELEFAX: (810) 539-5055
; INFORMATION FOR SEQ ID NO: 79:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-629-001A-79

Query Match 14.4%; Score 13; DB 2; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.9e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0;

QY 5 aaaaatttttc 17
|||||
DB 15 AAAAAATTAATTC 27

RESULT 3

US-08-642-274D-158
; Sequence 158, Application US/08642274D
; Patent No. 6200749
; GENERAL INFORMATION:
; APPLICANT: Shiloh, Yosef
; TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
; TITLE OF INVENTION: SCREEN FOR A PARTIAL A-T PHENOTYPE
; FILE REFERENCE: 229000033
; CURRENT APPLICATION NUMBER: US/08/642,274D
; CURRENT FILING DATE: 1996-05-03
; NUMBER OF SEQ ID NOS: 220
; SOFTWARE: Patent In Ver. 2.1
; SEQ ID NO 158

; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: intronic
US-08-642-274D-158

Query Match 14.4%; Score 13; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.9e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0;

QY 5 aaaaatttttc 17
|||||
DB 15 aaaaatttttc 27

RESULT 4

PCT-US96-00547-40/c
; Sequence 40, Application PC/TUS9600547
; GENERAL INFORMATION:
; APPLICANT: Virogenetics Corporation
; TITLE OF INVENTION: RECOMBINANT POXYVIRUS-HTML, COMPOSITIONS
; NUMBER OF SEQUENCES: 56
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Curtis, Morris & Safford, P.C.
; STREET: 530 Fifth Avenue, 25th Floor
; CITY: New York
; STATE: New York
; COUNTRY: U.S.A.
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US96/00547
; FILING DATE: 12-JAN-1996
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/372,664
; FILING DATE: 13-JAN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Frommer, William S.
; REGISTRATION NUMBER: 25,506
; REFERENCE/DOCKET NUMBER: 454310-2621
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 840-3333
; TELEFAX: (212) 840-0712
; INFORMATION FOR SEQ ID NO: 40:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 38 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
PCT-US96-00547-40

Query Match 14.4%; Score 13; DB 5; Length 38;
Best Local Similarity 100.0%; Pred. No. 6.8e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0;

QY 26 cgttaattttat 38
|||||
DB 25 CGTGATTTTAT 13

RESULT 5

US-08-105-483-168/c

Sequence 168, Application US/08105483
Patent No. 5494807
GENERAL INFORMATION:
APPLICANT: Paoletti, Enzo
TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
NUMBER OF SEQUENCES: 462
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESS: c/o William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/105,483
FILING DATE: 12-AUG-1993
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/847,951
FILING DATE: 06-MAR-1992
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2400
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-105-483-168

Query Match 14.4%; Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 26 ctgtaattttat 38
Db 25 CTGTAATTTTAT 13

RESULT 6
US-08-709-209-168/c
Sequence 168, Application US/08709209
Patent No. 5762938
GENERAL INFORMATION:
APPLICANT: Paoletti, Enzo
TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
NUMBER OF SEQUENCES: 462
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESS: c/o William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/105,483
FILING DATE: 12-AUG-1993
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/847,951
FILING DATE: 06-MAR-1992
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2400
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-709-209-168

SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/709,209
FILING DATE: 21-AUG-1996
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/105,483
FILING DATE: 12-AUG-1993
APPLICATION NUMBER: US 07/847,951
FILING DATE: 06-MAR-1992
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2400
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-709-209-168

Query Match 14.4%; Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 26 ctgtaattttat 38
Db 25 CTGTAATTTTAT 13

RESULT 7
US-08-303-275-56/c
Sequence 56, Application US/08303275
Patent No. 576598
GENERAL INFORMATION:
APPLICANT: Paoletti, Enzo
APPLICANT: Tartaglia, James
TITLE OF INVENTION: IMMUNODEFICIENCY VIRUS RECOMBINANT
NUMBER OF SEQUENCES: 205
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESS: c/o William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/303,275
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/897,382
FILING DATE: 11-JUN-1992
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2420
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712

INFORMATION FOR SEQ ID NO: 56;
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-303-275-56

Query Match 14.4%; Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. NO. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 cgtgaattttat 38
DB 25 CCGTAAATTTTAT 13

RESULT 8

US-08-458-101-168/c
Sequence 168, Application US/08458101

Patent No. 5766599

GENERAL INFORMATION:

APPLICANT: Paolucci, Enzo

APPLICANT: Perkus, Marion E.

APPLICANT: Taylor, Jill

APPLICANT: Tartaglia, James

APPLICANT: No. 5766599, Elizabeth K.

APPLICANT: Riviere, Michel

APPLICANT: de Taisne, Charles

APPLICANT: Limbach, Keith J.

APPLICANT: Johnson, Gerard P.

APPLICANT: Pincus, Steven E.

APPLICANT: Cox, William I.

APPLICANT: Audonnet, Jean-Christophe Francis

APPLICANT: Getlig, Russell Robert

TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE

TITLE OF INVENTION: STRAIN

NUMBER OF SEQUENCES: 467

CORRESPONDENCE ADDRESS:

ADDRESSEE: Curtis, Morris & Safford

ADDRESSEE: C/O William S. Frommer

STREET: 530 Fifth Avenue

CITY: New York

STATE: NY

COUNTRY: USA

ZIP: 10036

COMPUTER READABLE FORM:

MEDIUM TYPE: floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/458,101

FILING DATE: 01-JUN-1995

CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:

NAME: Frommer, William S.

REGISTRATION NUMBER: 25,506

REFERENCE/DOCKET NUMBER: 45310-2740

TELEPHONE: (212) 840-3333

TELEPHONE: (212) 840-0712

INFORMATION FOR SEQ ID NO: 168:

SEQUENCE CHARACTERISTICS:

LENGTH: 39 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

US-08-458-101-168

Query Match 14.4%; Score 13; DB 1; Length 39;

Best Local Similarity 100.0%; Pred. No. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 cgtgaattttat 38
DB 25 CCGTAAATTTTAT 13

RESULT 9

US-08-271-880A-201

Sequence 201, Application US/08271880A

Patent No. 5693535

GENERAL INFORMATION:

APPLICANT: Kenneth G. Draper

APPLICANT: Bharat Chowrira

APPLICANT: James McSwigen

APPLICANT: Dan T. Stinchcomb

APPLICANT: James D. Thompson

TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING

TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS

NUMBER OF SEQUENCES: 232

CORRESPONDENCE ADDRESS:

ADDRESSEE: Lyon & Lyon

STREET: 633 West Fifth Street

STREET: Suite 4700

CITY: Los Angeles

STATE: California

COUNTRY: U.S.A.

ZIP: 90071

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5" Diskette, 1.44 MB

MEDIUM TYPE: Storage

COMPUTER: IBM Compatible

OPERATING SYSTEM: IBM P.C. DOS 5.0

SOFTWARE: FASTSEQ Version 1.5

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/271,880A

FILING DATE: July 7, 1994

PRIOR APPLICATION DATA:

PRIOR APPLICATION DATA: including Application

PRIOR APPLICATION DATA: described below:

APPLICATION NUMBER: 08/103,243

FILING DATE: August 6, 1993

APPLICATION NUMBER: 07/882,886

FILING DATE: May 14, 1992

ATTORNEY/AGENT INFORMATION:

NAME: Waidburg, Richard

REGISTRATION NUMBER: 32,327

REFERENCE/DOCKET NUMBER: 206/116

TELECOMMUNICATION INFORMATION:

TELEPHONE: (213) 489-1600

TELEFAX: (213) 955-0440

TELEX: 67-3510

INFORMATION FOR SEQ ID NO: 201:

SEQUENCE CHARACTERISTICS:

LENGTH: 14 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

US-08-271-880A-201

Query Match 13.3%; Score 12; DB 1; Length 14;
Best Local Similarity 100.0%; Pred. NO. 2.2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
DB 2 CAATGAAGCAA 13

RESULT 10

US-08-910-408-201
Sequence 201, Application US/08910408
Patent No. 5972704
GENERAL INFORMATION:
APPLICANT: Kenneth G. Diaper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITTING
TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910,408
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/271,880
FILING DATE: July 7, 1994
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 201:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-910-408-201

Query Match 13.3%; Score 12; DB 2; Length 14;
Best Local Similarity 100.0%; Pred. No. 2.2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagcaaa 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 11
US-09-249-215-201
Sequence 201, Application US/09249215
Patent No. 6159692
GENERAL INFORMATION:
APPLICANT: Kenneth G. Diaper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson

TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITTING
HUMAN IMMUNODEFICIENCY VIRUS
REPLICATION
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/249,215
FILING DATE: 12-Feb-1999
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/910,408
FILING DATE: <Unknown>
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 201:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 201:
US-09-249-215-201

Query Match 13.3%; Score 12; DB 3; Length 14;
Best Local Similarity 100.0%; Pred. No. 2.2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagcaaa 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 12
US-09-163-162-19
Sequence 19, Application US/09163162
Patent No. 6077709
GENERAL INFORMATION:
APPLICANT: Bennett, C. Frank
APPLICANT: Ackermann, Elizabeth J.
APPLICANT: Swayze, Eric E.
APPLICANT: Cowsett, Lex M.
TITLE OF INVENTION: ANTISENSE MODULATION OF SURVIVIN EXPRESSION
FILE REFERENCE: RTS-0008
CURRENT APPLICATION NUMBER: US/09/163,162
CURRENT FILING DATE: 1998-09-29
NUMBER OF SEQ ID NOS: 47
SEQ ID NO 19
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence

FEATURE:
OTHER INFORMATION: Antisense Oligonucleotide
US-09-163-162-19

Query Match 13.3%; Score 12; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 51 tcttatgttg 62
Db 1 tcttatgttg 12

RESULT 13
US-09-286-407-19
Sequence 19, Application US/09286407A
Patent No. 6165788
GENERAL INFORMATION:
APPLICANT: Bennett, C. Frank
APPLICANT: Ackermann, Elizabeth J.
APPLICANT: Swayze, Eric E.
APPLICANT: Cowser, Lex M.
TITLE OF INVENTION: ANTISENSE MODULATION OF SURVIVIN EXPRESSION
FILE REFERENCE: ISPH-0349
CURRENT APPLICATION NUMBER: US/09/286,407A
CURRENT FILING DATE: 1999-04-05
NUMBER OF SEQ ID NOS: 48
SEQ ID NO 19
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Antisense Oligonucleotide
US-09-286-407-19

Query Match 13.3%; Score 12; DB 4; Length 18;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 51 tcttatgttg 62
Db 1 tcttatgttg 12

RESULT 14
US-08-661-507-6/C
Sequence 6, Application US/08661507
Patent No. 5814490
GENERAL INFORMATION:
APPLICANT: Spears, Patricia A.
TITLE OF INVENTION: AMPLIFICATION AND DETECTION OF CHLAMYDIA
TITLE OF INVENTION: TRACHOMATIS NUCLEIC ACIDS
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
STREET: 1 Becton Drive
CITY: Franklin Lakes
STATE: NJ
COUNTRY: US
ZIP: 07417
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/661,507
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Fugit, Donna R.

REGISTRATION NUMBER: 32,135
REFERENCE/DOCKET NUMBER: P-3489
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-661-507-6

Query Match 13.3%; Score 12; DB 1; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcacgttt 50
Db 13 gtttcacgttt 2

RESULT 15
US-08-855-085-4/C
Sequence 4, Application US/08855085
Patent No. 5846726
GENERAL INFORMATION:
APPLICANT: Nadeau, James G.
APPLICANT: Pitner, James B.
APPLICANT: Schram, James L.
APPLICANT: Linn, Carl P.
APPLICANT: Vonk, Glenn P.
APPLICANT: Walker, George T.
TITLE OF INVENTION: Detection of Nucleic Acids by
TITLE OF INVENTION: Fluorescence Quenching
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
STREET: 1 Becton Drive
CITY: Franklin Lakes
STATE: NJ
COUNTRY: US
ZIP: 07417
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/855,085
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Fugit, Donna R.
REGISTRATION NUMBER: 32,135
REFERENCE/DOCKET NUMBER: P-3747
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-855-085-4

Query Match 13.3%; Score 12; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcacgttt 50
Db 13 gtttcacgttt 2

RESULT 16

US-09-186-030-4/c
; Sequence 4, Application US/09186030
; Patent No. 5919630
; GENERAL INFORMATION:
; APPLICANT: Nadeau, James G.
; APPLICANT: Pitner, James B.
; APPLICANT: Schram, James L.
; APPLICANT: Linn, Carl P.
; APPLICANT: Vonk, Glenn P.
; APPLICANT: Walker, George T.
; TITLE OF INVENTION: Detection of Nucleic Acids by
; TITLE OF INVENTION: Fluorescence Quenching
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
; STREET: 1 Becton Drive
; CITY: Franklin Lakes
; STATE: NJ
; COUNTRY: US
; ZIP: 07417
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/186,030
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/855,085
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Fugit, Donna R.
; REGISTRATION NUMBER: 32,135
; REFERENCE/DOCKET NUMBER: P-3747
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-09-186-030-4

Query Match 13.3%, Score 12; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
DB 13 GTTTCACGTTT 2

RESULT 17
US-08-865-675-4/c
; Sequence 4, Application US/08865675
; Patent No. 5928869
; GENERAL INFORMATION:
; APPLICANT: Nadeau, James G.
; APPLICANT: Pitner, James B.
; APPLICANT: Linn, Carl P.
; APPLICANT: Schram, James L.
; APPLICANT: Vonk, Glenn P.
; APPLICANT: Walker, George T.
; TITLE OF INVENTION: Detection of Nucleic Acids by
; TITLE OF INVENTION: FLUORESCENCE QUENCHING
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
; STREET: 1 Becton Drive
; CITY: Franklin Lakes
; STATE: NJ
; COUNTRY: US
; ZIP: 07417

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/865,675
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Fugit, Donna R.
; REGISTRATION NUMBER: 32,135
; REFERENCE/DOCKET NUMBER: P-3746
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-865-675-4

Query Match 13.3%, Score 12; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
DB 13 GTTTCACGTTT 2

RESULT 18
US-08-933-749-5/c
; Sequence 5, Application US/08933749
; Patent No. 5935791
; GENERAL INFORMATION:
; APPLICANT: Nadeau, James G.
; APPLICANT: Hsieh, Helen V.
; APPLICANT: Pitner, James B.
; APPLICANT: Linn, Carl P.
; TITLE OF INVENTION: Detection of Nucleic Acids by
; TITLE OF INVENTION: Fluorescence Quenching
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
; STREET: 1 Becton Drive
; CITY: Franklin Lakes
; STATE: NJ
; COUNTRY: US
; ZIP: 07417
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/933,749
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Fugit, Donna R.
; REGISTRATION NUMBER: 32,135
; REFERENCE/DOCKET NUMBER: P-3749
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-933-749-5

Query Match 13.3%, Score 12; DB 2; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATCTTT 2

RESULT 19
US-09-237-510-4/C

; Sequence 4, Application US/09237510
; Patent No. 5958700

; GENERAL INFORMATION:

; APPLICANT: Nadeau, James G.

; APPLICANT: Pitner, James B.

; APPLICANT: Linn, Carl P.

; APPLICANT: Schram, James L.

; TITLE OF INVENTION: DETECTION OF NUCLEIC ACIDS BY

; NUMBER OF SEQUENCES: 9

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company

; STREET: 1 Becton Drive

; CITY: Franklin Lakes

; STATE: NJ

; COUNTRY: US

; ZIP: 07417

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/237,510

; FILING DATE:

; CLASSIFICATION:

; ATTORNEY/AGENT INFORMATION:

; NAME: Fugit, Donna R.

; REGISTRATION NUMBER: 32,135

; REFERENCE/DOCKET NUMBER: P-3746

; INFORMATION FOR SEQ ID NO: 4:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 21 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; US-09-237-510-4

; Query Match

; Best Local Similarity 13.3%; Score 12; DB 2; Length 21;

; Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATCTTT 2

RESULT 20
US-09-120-916-4/C

; Sequence 4, Application US/09120916

; Patent No. 6054279

; GENERAL INFORMATION:

; APPLICANT: Nadeau, James G.

; APPLICANT: Pitner, James B.

; APPLICANT: Schram, James L.

; APPLICANT: Linn, Carl P.

; APPLICANT: Vonk, Glenn P.

; APPLICANT: Walker, George T.

; TITLE OF INVENTION: Detection of Nucleic Acids by

; NUMBER OF SEQUENCES: 6

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
; STREET: 1 Becton Drive
; CITY: Franklin Lakes
; STATE: NJ
; COUNTRY: US

; ZIP: 07417

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/120,916

; FILING DATE:

; CLASSIFICATION:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/855,085

; FILING DATE:

; ATTORNEY/AGENT INFORMATION:

; NAME: Fugit, Donna R.

; REGISTRATION NUMBER: 32,135

; REFERENCE/DOCKET NUMBER: P-3747

; INFORMATION FOR SEQ ID NO: 4:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 21 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; US-09-120-916-4

; Query Match

; Best Local Similarity 13.3%; Score 12; DB 3; Length 21;

; Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATCTTT 2

RESULT 21
US-08-964-020-8/C

; Sequence 8, Application US/08964020

; Patent No. 6077669

; GENERAL INFORMATION:

; APPLICANT: Vonk, Glenn P.

; APPLICANT: Little, Michael C.

; TITLE OF INVENTION: Kit and Method for Fluorescence Based

; NUMBER OF SEQUENCES: 20

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Richard J. Rodrick - Becton, Dickinson and

; ADDRESS: Company

; STREET: 1 Becton Drive

; CITY: Franklin Lakes

; STATE: NJ

; COUNTRY: USA

; ZIP: 07417

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/964,020

; FILING DATE:

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Higbet, David W.

; REGISTRATION NUMBER: 30,265

; REFERENCE/DOCKET NUMBER: P-4025

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (201) 847-5317

TELEFAX: (201) 848-9228
: INFORMATION FOR SEQ ID NO: 8:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 21 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-08-964-020-8

Query Match 13.3%; Score 12; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
DB 13 GTTTCACGTTT 2

RESULT 22
US-09-235-583-5/c
: Sequence 5, Application US/09235583
: Patent No. 6130047
: GENERAL INFORMATION:
: APPLICANT: Nadeau, James G.
: APPLICANT: Hsieh, Helen V.
: APPLICANT: Pitner, James B.
: APPLICANT: Linn, Carl P.
: TITLE OF INVENTION: Detection of Nucleic Acids by
: NUMBER OF SEQUENCES: 10
: CORRESPONDENCE ADDRESSES:
: ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
: STREET: 1 Becton Drive
: CITY: Franklin Lakes
: STATE: NJ
: COUNTRY: US
: ZIP: 07417
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/235,583
: FILING DATE:
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Fugit, Donna R.
: REGISTRATION NUMBER: 32,135
: REFERENCE/DOCKET NUMBER: P-3749
: INFORMATION FOR SEQ ID NO: 5:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 21 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-09-235-583-5

Query Match 13.3%; Score 12; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
DB 13 GTTTCACGTTT 2

RESULT 23
US-09-599-164-5/c
: Sequence 5, Application US/09599164
: Patent No. 6261784

GENERAL INFORMATION:
: APPLICANT: Nadeau, James G.
: APPLICANT: Hsieh, Helen V.
: APPLICANT: Pitner, James B.
: APPLICANT: Linn, Carl P.
: TITLE OF INVENTION: Detection of Nucleic Acids by
: NUMBER OF SEQUENCES: 10
: CORRESPONDENCE ADDRESSES:
: ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
: STREET: 1 Becton Drive
: CITY: Franklin Lakes
: STATE: NJ
: COUNTRY: US
: ZIP: 07417

COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/599,164
: FILING DATE:
: CLASSIFICATION:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US/08/933,749
: FILING DATE:
: ATTORNEY/AGENT INFORMATION:
: NAME: Fugit, Donna R.
: REGISTRATION NUMBER: 32,135
: REFERENCE/DOCKET NUMBER: P-3749
: INFORMATION FOR SEQ ID NO: 5:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 21 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-09-599-164-5

Query Match 13.3%; Score 12; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
DB 13 GTTTCACGTTT 2

RESULT 24
US-08-727-003A-10/c
: Sequence 10, Application US/08727003A
: Patent No. 5804383
: GENERAL INFORMATION:
: APPLICANT: Gruenert, Dieter, C.
: APPLICANT: Dohman, Austin F.
: TITLE OF INVENTION: A METHOD AND ASSAY FOR
: TITLE OF INVENTION: DETECTION OF THE EXPRESSION
: TITLE OF INVENTION: OF ALLELE-SPECIFIC MUTATIONS
: TITLE OF INVENTION: BY ALLELE-SPECIFIC IN SITU
: TITLE OF INVENTION: REVERSE TRANSCRIPTASE
: TITLE OF INVENTION: POLYMERASE CHAIN REACTION
: NUMBER OF SEQUENCES: 55
: CORRESPONDENCE ADDRESSES:
: ADDRESSEE: PETERS, VERNY, JONES & BIK A, L.L.P.
: STREET: 385 Sherman Avenue
: CITY: Palo Alto
: STATE: California
: COUNTRY: United States of America
: ZIP: 94306-1840
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Diskette - 3.5 inch, 1.44 Kb storage
: COMPUTER: PC

OPERATING SYSTEM: DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA: 003A
APPLICATION NUMBER: US/08/727,003A
FILING DATE: October 8, 1996
CLASSIFICATION: 435
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 60/005,254
FILING DATE: October 10, 1995
ATTORNEY/AGENT INFORMATION:
NAME: Hana Verry
REGISTRATION NUMBER: 30,518
REFERENCE/DOCKET NUMBER: 480-77
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415)324-1677
TELEFAX: (415)324-1678
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 23 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: synthetic oligonucleotide
US-08-727-003A-10

Query Match 13.3%; Score 12; DB 1; Length 23;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0;

Oy 59 ttggagcaataa 70
|||||
Db 23 TTGGAGCAATAA 12

RESULT 25
US-08-487-799-11/c
Sequence 11, Application US/08487799C
Patent No. 6010908
GENERAL INFORMATION:
APPLICANT: Gruenert, Dieter C.
APPLICANT: Kunzelmann, Karl
TITLE OF INVENTION: GENE THERAPY BY SMALL FRAGMENTS HOMOLOGOUS REPLACEMENT
FILE REFERENCE: 480.18-1(HV)
CURRENT APPLICATION NUMBER: US/08/487,799C
CURRENT FILING DATE: 1995-06-07
EARLIER APPLICATION NUMBER: 07/933,471
EARLIER FILING DATE: 1992-08-21
EARLIER APPLICATION NUMBER: 08/409,544
EARLIER FILING DATE: 1995-03-24
NUMBER OF SEQ ID NOS: 87
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 11
LENGTH: 23
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: synthetic
US-08-487-799-11

Query Match 13.3%; Score 12; DB 3; Length 23;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0;

Oy 59 ttggagcaataa 70
|||||
Db 23 TTGGAGCAATAA 12

RESULT 26
US-08-210-762E-31

Sequence 31, Application US/08210762E
Patent No. 5837441
GENERAL INFORMATION:
APPLICANT: Hjelle, Brian
APPLICANT: Jensen, Steve
TITLE OF INVENTION: Molecular Clones Producing Recombinant DNA Antigens of
TITLE OF INVENTION: the HARDS Virus.
NUMBER OF SEQUENCES: 85
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hoffman, Wasson & Gittler
STREET: 2361 Jefferson Davis Highway
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 500 Kb storage
COMPUTER: Accel 486
OPERATING SYSTEM: Windows 3.1
SOFTWARE: Wordperfect 6.1 for Windows
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/210,762E
FILING DATE: 22-MAR-94
CLASSIFICATION: 435
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/141,035
FILING DATE: 26-OCT-93
APPLICATION NUMBER: 08/120,096
FILING DATE: 13-SEP-93
APPLICATION NUMBER: 08/111,519
FILING DATE: 25-AUG-93
ATTORNEY/AGENT INFORMATION:
NAME: Butml, Jean A.
REGISTRATION NUMBER: 24,236
REFERENCE/DOCKET NUMBER: A4710CIP3.SL3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)415-0100
TELEFAX: (703)418-2768
INFORMATION FOR SEQ ID NO: 31:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA viral
HYPOTHETICAL: no
ANTI-SENSE: no
ORIGINAL SOURCE:
ORGANISM: Four Corners Hantavirus
INDIVIDUAL ISOLATE: 3H226
IMMEDIATE SOURCE:
LIBRARY:
CLONE:
PUBLICATION INFORMATION:
AUTHORS: Hjelle, Brian
AUTHORS: Jensen, Steven
AUTHORS: Torrez-Martinez, No. 5837441ah
AUTHORS: Yamada, Takashi
AUTHORS: No. 5837441te, Kurt
AUTHORS: Zumwalt, Ross
AUTHORS: MacInnes, Kersti
AUTHORS: Myers, Gerald
TITLE: A No. 5837441el Hantavirus Associated with an Outbreak of Fatal Respir
TITLE: Disease in the Southwestern United States: Evolutionary Relationships
TITLE: Hantaviruses-Running Title: Hantavirus-associated ARDS
JOURNAL: Journal of Virology
VOLUME: 68
PAGES: in press
DATE: 1994
RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 24
US-08-210-762E-31

Query Match 13.3%; Score 12; DB 2; Length 24;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 21 ttatctgtaatt 32
|||||
Db 2 GTTACTGTAATT 13

RESULT 27
US-08-256-799-26
Sequence 26, Application US/08256799
Patent No. 6222094

GENERAL INFORMATION:
APPLICANT: HANSSON, Lennart
APPLICANT: STROEMOYIST, Mats
APPLICANT: BERGSTROM, Sven
APPLICANT: HERNNELL, Olle
APPLICANT: Toernell, Jan
TITLE OF INVENTION: DNA ENCODING KAPPA-CASEIN, PROCESS FOR
OBTAINING THE PROTEIN AND USE THEREOF
NUMBER OF SEQUENCES: 26
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWDY AND NEIMARK
STREET: 419 Seventh Street, N.W., Suite 300
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20004

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/256,799
FILING DATE: 06-DEC-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DK 88/92
FILING DATE: 23-JAN-1992
ATTORNEY/AGENT INFORMATION:
NAME: COOPER, Iver P.
REGISTRATION NUMBER: 28,005
REFERENCE/DOCKET NUMBER: HANSSON-1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-737-3528
TELEFAX: 202-737-3528
INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-256-799-26

Query Match 13.3%; Score 12; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 35 ttatgtttcat 46
|||||
Db 12 TTATGTTTCAT 23

RESULT 28
US-08-462-437-26
Sequence 26, Application US/08462437
Patent No. 6232094
GENERAL INFORMATION:
APPLICANT: HANSSON, Lennart

APPLICANT: STROEMOYIST, Mats
APPLICANT: BERGSTROM, Sven
APPLICANT: HERNNELL, Olle
APPLICANT: Toernell, Jan
TITLE OF INVENTION: DNA ENCODING KAPPA-CASEIN, PROCESS
OBTAINING THE PROTEIN AND USE THEREOF
NUMBER OF SEQUENCES: 31
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWDY AND NEIMARK
STREET: 419 Seventh Street, N.W., Suite 300
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20004

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/462,437
FILING DATE: 05-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DK 88/92
FILING DATE: 23-JAN-1992
ATTORNEY/AGENT INFORMATION:
NAME: COOPER, Iver P.
REGISTRATION NUMBER: 28,005
REFERENCE/DOCKET NUMBER: HANSSON-1A
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-628-5197
TELEFAX: 202-737-3528
TELEX: 248633

INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-462-437-26

Query Match 13.3%; Score 12; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 35 ttatgtttcat 46
|||||
Db 12 TTATGTTTCAT 23

RESULT 29
US-08-271-880A-140/C
Sequence 140, Application US/08271880A
Patent No. 5693535

GENERAL INFORMATION:
APPLICANT: Kenneth G. Draper
APPLICANT: Bharat Chowrira
APPLICANT: James McSwigen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING
HUMAN IMMUNODEFICIENCY VIRUS
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.

ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/271,880A
FILING DATE: July 7, 1994
PRIOR APPLICATION DATA:
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 140:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-271-880A-140

Query Match 13.3%; Score 12; DB 1; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 caatgaagca 85
Db 23 CAATGAAGCAA 12

RESULT 30
US-08-910-408-140/C
Sequence 140, Application US/08910408
Patent No. 59/2704
GENERAL INFORMATION:
APPLICANT: Kenneth G. Draper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING
TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS
REPLICATION
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910,408

FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/271,880
FILING DATE: July 7, 1994
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 140:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-910-408-140

Query Match 13.3%; Score 12; DB 2; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 caatgaagca 85
Db 23 CAATGAAGCAA 12

RESULT 31
US-09-249-215-140/C
Sequence 140, Application US/09249215
Patent No. 6159692
GENERAL INFORMATION:
APPLICANT: Kenneth G. Draper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING
TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS
REPLICATION
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/249,215
FILING DATE: 12-Feb-1999
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/910,408
FILING DATE: <unknown>
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:

NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 140:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 140:
US-09-249-215-140

Query Match 13.3%; Score 12; DB 3; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 74 caatgaagcaa 85
|||
Db 23 CAATGAAGCAA 12

RESULT 32
US-08-835-728D-72
Sequence 72, Application US/08835728D
Patent No. 6017704
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylin, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/835,728D
FILING DATE: April 11, 1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/656,716
FILING DATE: June 03, 1996,
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5099
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 72:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-835-728D-72

Query Match 13.3%; Score 12; DB 3; Length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 32 ttttaagtgtt 43
|||||
Db 6 TTTTATGTTTT 17

RESULT 33
US-08-835-728D-176/C
Sequence 176, Application US/08835728D
Patent No. 6017704
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylin, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/835,728D
FILING DATE: April 11, 1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/656,716
FILING DATE: June 03, 1996,
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5099
TELEFAX: 619/678-5070
INFORMATION FOR SEQ ID NO: 176:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-835-728D-176

Query Match 13.3%; Score 12; DB 3; Length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 32 ttttaagtgtt 43
|||||
Db 21 TTTTATGTTTT 10

RESULT 34
US-09-490-558-72
Sequence 72, Application US/09490558
Patent No. 6265171
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylin, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400

CITY: LA JOLLA
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/490,558
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/835,728
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 72:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-09-490-558-72

Query Match 13.3% Score 12; DB 4; length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 32 ttttatgttt 43
|||||
DB 6 ttttatgttt 17

RESULT 35
US-09-490-558-176/C
Sequence 176, Application US/09490558
Patent No. 6265171
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylin, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSER: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: LA JOLLA
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/490,558
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/835,728
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347

REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 176:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-09-490-558-176

Query Match 13.3% Score 12; DB 4; length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 32 ttttatgttt 43
|||||
DB 21 ttttatgttt 10

RESULT 36
US-08-668-128B-6/C
Sequence 6, Application US/08668128B
Patent No. 5840568
GENERAL INFORMATION:
APPLICANT: Fireundschuh, Michael
TITLE OF INVENTION: Hodgkin's Disease Associated Molecules And
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSER: Felte & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/668,128B
FILING DATE: 21-JUNE-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/644,116
FILING DATE: 10-MAY-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/580,980
FILING DATE: 03-JANUARY-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/479,328
FILING DATE: 07-JUNE-1995
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 5840568man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5441
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 27 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-668-128B-6

Query Match 13.3% Score 12; DB 2; length 27;

Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagt 88
|||||
Db 26 TGAAGCAAGTG 15

RESULT 37
US-08-905-445-6/c
; Sequence 6, Application US/08905445
; Patent No. 5864015
; GENERAL INFORMATION:
; APPLICANT: Pfeundschnub, Michael
; TITLE OF INVENTION: Hodgkin's Disease Associated Molecules And
; TITLE OF INVENTION: Uses Thereof
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Felfe & Lynch
; STREET: 805 Third Avenue
; CITY: New York City
; STATE: New York
; ZIP: 10022
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
; COMPUTER: IBM
; OPERATING SYSTEM: PC-DOS
; SOFTWARE: Wordperfect
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/905,445
; FILING DATE: 04-AUG-1997
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/668,128
; FILING DATE: 21-JUNE-1996
; APPLICATION NUMBER: 08/644,116
; FILING DATE: 10-MAY-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/580,980
; FILING DATE: 03-JANUARY-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/479,328
; FILING DATE: 07-JUNE-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Hanson, No. 5864015man D.
; REGISTRATION NUMBER: 30,946
; REFERENCE/DOCKET NUMBER: LUD 5441
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 688-9200
; TELEFAX: (212) 838-3884
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-905-445-6

Query Match 13.3%; Score 12; DB 2; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagt 88
|||||
Db 26 TGAAGCAAGTG 15

RESULT 38
US-08-644-116A-6/c
; Sequence 6, Application US/08644116A
; Patent No. 6140464
; GENERAL INFORMATION:

APPLICANT: Pfeundschnub, Michael; Rammensee, Hans-Georg
TITLE OF INVENTION: Method For Identifying Or Isolating A Molecule
TITLE OF INVENTION: And Molecules Identified Thereby
NUMBER OF SEQUENCES: 13
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/644,116A
FILING DATE: 10-MAY-1996
CLASSIFICATION: 436
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/580,980
FILING DATE: 03-JANUARY-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/479,328
FILING DATE: 07-JUNE-1995
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 6140464man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5410.2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 27 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-644-116A-6

Query Match 13.3%; Score 12; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagt 88
|||||
Db 26 TGAAGCAAGTG 15

RESULT 39
US-08-771-602D-20
; Sequence 20, Application US/08771602D
; Patent No. 5976795
; GENERAL INFORMATION:
; APPLICANT: Voytas, Daniel F.
; APPLICANT: Zou, Siye
; TITLE OF INVENTION: Retrotransposon and Methods
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Greenlee, Winner and Sullivan, P.C.
; STREET: 5370 Manhattan Circle, Suite 201
; CITY: Boulder
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80303
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/771,602D

FILING DATE: 20-DEC-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/010,869
FILING DATE: 31-JAN-1996
ATTORNEY/AGENT INFORMATION:
NAME: Farber, Donna M.
REGISTRATION NUMBER: 33,878
REFERENCE/DOCKET NUMBER: 8-96
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 499-8080
TELEFAX: (303) 499-8089
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 28 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "Oligonucleotide."
HYPOTHETICAL: NO
US-08-771-602D-20

Query Match 13.3%; Score 12; DB 2; Length 28;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 31 attttaattt 42
|||||
DB 7 ATTATATGTTT 18

RESULT 40
US-08-771-602D-21/C
Sequence 21, Application US/08771602D
Patent No. 5976795
GENERAL INFORMATION:
APPLICANT: Voytas, Daniel F.
TITLE OF INVENTION: Retrotransposon and Methods
NUMBER OF SEQUENCES: 51
CORRESPONDENCE ADDRESS:
ADDRESSEE: Greenlee, Winner and Sullivan, P.C.
STREET: 5370 Manhattan Circle, Suite 201
CITY: Boulder
STATE: Colorado
COUNTRY: USA
ZIP: 80303
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/771,602D
FILING DATE: 20-DEC-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/010,869
FILING DATE: 31-JAN-1996
ATTORNEY/AGENT INFORMATION:
NAME: Farber, Donna M.
REGISTRATION NUMBER: 33,878
REFERENCE/DOCKET NUMBER: 8-96
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 499-8080
TELEFAX: (303) 499-8089
INFORMATION FOR SEQ ID NO: 21:
SEQUENCE CHARACTERISTICS:
LENGTH: 28 base pairs
TYPE: nucleic acid
STRANDEDNESS: single

TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "Oligonucleotide."
HYPOTHETICAL: NO
US-08-771-602D-21

Query Match 13.3%; Score 12; DB 2; Length 28;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 31 attttaattt 42
|||||
DB 22 ATTATATGTTT 11

RESULT 41
US-08-619-724-3/C
Sequence 3, Application US/08619724
Patent No. 5827653
GENERAL INFORMATION:
APPLICANT: SAMMES, Peter George
APPLICANT: GARMAN, Andrew John
TITLE OF INVENTION: NUCLEIC ACID DETECTION WITH ENERGY TRANSFER
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: PILLSBURY MADISON & SUTRO, L.L.P.
STREET: 1100 New York Avenue, N.W.
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20005-3918
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Microsoft Word
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/619,724
FILING DATE: 20-MAY-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/GB94/02068
FILING DATE: 23-SEP-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9412106.8
FILING DATE: 16-JUN-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9319826.5
FILING DATE: 23-SEP-1993
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
US-08-619-724-3

Query Match 13.3%; Score 12; DB 1; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 cgttaatttta 37
|||||
DB 29 CTTATATTTTA 18

RESULT 42
US-08-629-001A-25/C
Sequence 25, Application US/08629001A
Patent No. 5858661
GENERAL INFORMATION:

APPLICANT: Shiloh, Yosef
TITLE OF INVENTION: ATAXIA-TELANGIECTASIA GENE AND ITS
NUMBER OF INVENTION: GENOMIC ORGANIZATION
NUMBER OF SEQUENCES: 139
CORRESPONDENCE ADDRESS:
ADDRESSEE: Kohn & Associates
STREET: 30500 No. 5858661thwestern Hwy.
CITY: Farmington Hills
STATE: Michigan
COUNTRY: US
ZIP: 48334
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/629,001A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Kohn, Kenneth I.
REGISTRATION NUMBER: 30,955
REFERENCE/DOCKET NUMBER: 2290,00032
TELECOMMUNICATION INFORMATION:
TELEPHONE: (810) 539-5050
TELEFAX: (810) 539-5055
INFORMATION FOR SEQ ID NO: 25:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-629-001A-25

Query Match 13.3%; Score 12; DB 2; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 aaaaaattatt 15
Db 19 AAAAAATTATT 8

RESULT 43
US-08-642-274D-104/c
; Sequence 104, Application US/08642274D
; Patent No. 6200749
; GENERAL INFORMATION:
; APPLICANT: Shiloh, Yosef
; TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
; FILE REFERENCE: 229000033
; CURRENT APPLICATION NUMBER: US/08/642,274D
; CURRENT FILING DATE: 1996-05-03
; NUMBER OF SEQ ID NOS: 220
; SOFTWARE: Patentln Ver. 2.1
; SEQ ID NO 104
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence:intronic
; OTHER INFORMATION: sequence
US-08-642-274D-104

Query Match 13.3%; Score 12; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 4 aaaaaattatt 15

Db 19 AAAAAATTATT 8

RESULT 44
US-08-935-312-14/c
; Sequence 14, Application US/08935312
; Patent No. 6207455
; GENERAL INFORMATION:
; APPLICANT: CHANG, Lung-Ji
; TITLE OF INVENTION: LENTIVIRAL VECTORS
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BROWDY AND NEIMARK, P.L.L.C.
; STREET: 624 Ninth Street, N.W.
; CITY: Washington
; STATE: D.C.
; COUNTRY: USA
; ZIP: 20001
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentln Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/935,312
; FILING DATE: 22-SEP-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: COOPER, Iver P.
; REGISTRATION NUMBER: 28,005
; REFERENCE/DOCKET NUMBER: CHANG-112
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-628-5197
; TELEFAX: 202-737-3528
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "DNA"
US-08-935-312-14

Query Match 13.3%; Score 12; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 74 caatgaagcaa 85
Db 28 CAATGAAGCAA 17

RESULT 45
US-08-341-456A-14/c
; Sequence 14, Application US/08341456A
; Patent No. 5767074
; GENERAL INFORMATION:
; APPLICANT: Besmer, Peter
; APPLICANT: NO. 5767074ka, Karl
; APPLICANT: Buck, Jochen
; APPLICANT: Moore, Malcolm A.S.
; TITLE OF INVENTION: COMPOSITIONS OF SOLUBLE C-KIT LIGAND AND
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Cooper & Dunham LLP
; STREET: 1185 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: U.S.A.

Query Match 13.3%; Score 12; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 74 caatgaagcaa 85
Db 28 CAATGAAGCAA 17

ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/341,456A
FILING DATE: 17-NOV-1994
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: White, John P.
REGISTRATION NUMBER: 28, 678
REFERENCE/DOCKET NUMBER: 37454-C1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 278-0400
TELEFAX: (212) 391-0525
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 33 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
US-08-341-456A-14

Query Match 13.3%; Score 12; DB 1; length 33;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 42 ttcatgtttct 53
|||||
Db 21 TTCATCTTTCT 10

Search completed: January 24, 2002, 03:24:26
Job time: 3713 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:57:29 ; Search time 2099.46 Seconds
(without alignments)
460.652 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
Sequence: 1 atgaaaaaattatttcaaa.....gtccaatgaagaacagtgcga 90

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 11351937 seqs, 5372889281 residues

Word size: 0

Total number of hits satisfying chosen parameters: 80718

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

EST:*
1: em_estfun:*
2: em_esthum:*
3: em_estlin:*
4: em_estom:*
5: em_estpl:*
6: em_estba:*
7: em_estro:*
8: em_estov:*
9: em_htc:*
10: gb_estl:*
11: gb_estl2:*
12: gb_htc:*
13: gb_gss:*
14: em_gss_fun:*
15: em_gss_hum:*
16: em_gss_inv:*
17: em_gss_pln:*
18: em_gss_pro:*
19: em_gss_rtd:*
20: em_gss_vrt:*
21: em_gss_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	14	15.6	50	13	TA154FI00
2	13	14.4	25	13	AZ829725
3	13	14.4	26	13	TA138120
4	13	14.4	28	13	AZ432653
5	13	14.4	36	13	AZ314238
6	13	14.4	49	10	AA500776
7	13	14.4	50	10	AU102750
8	12	13.3	23	13	AZ579583
9	12	13.3	24	13	AZ996858
10	12	13.3	25	13	AZ345616
11	12	13.3	27	10	AW246455
12	12	13.3	30	13	AZ610538

13	12	13.3	33	13	AZ507698
14	12	13.3	38	13	AZ487251
15	12	13.3	41	13	AZ662545
16	12	13.3	42	11	T17635
17	12	13.3	45	13	AZ514394
18	12	13.3	46	10	A1805932
19	12	13.3	48	13	AZ832106
20	12	13.3	50	11	D19972
21	11	12.2	20	13	AZ782616
22	11	12.2	21	13	AZ768088
23	11	12.2	22	10	AA991150
24	11	12.2	23	13	AZ766483
25	11	12.2	22	13	AZ387817
26	11	12.2	24	13	TA242F03P
27	11	12.2	25	13	AZ462654
28	11	12.2	28	13	AZ809415
29	11	12.2	29	11	D18726
30	11	12.2	29	13	AZ580321
31	11	12.2	31	13	AZ579378
32	11	12.2	32	13	AZ331642
33	11	12.2	32	13	AZ591923
34	11	12.2	33	11	U44209
35	11	12.2	33	13	AZ423204
36	11	12.2	35	10	AU014463
37	11	12.2	35	11	D19895
38	11	12.2	36	10	AU111149
39	11	12.2	36	13	AZ381596
40	11	12.2	36	13	AZ807406
41	11	12.2	37	10	AA647854
42	11	12.2	37	10	A1200438
43	11	12.2	37	13	AZ806836
44	11	12.2	38	11	D21038
45	11	12.2	38	11	D21038

ALIGNMENTS

RESULT 1
TA154FI00
LOCUS
DEFINITION
T. brucei sheared genomic DNA clone 154FI0, reverse sequence,
genomic survey sequence.

ACCESSION
AL473287
AL473287.1 GI:11838560

VERSION
GSS.

KEYWORDS
SOURCE
ORGANISM
Trypanosoma brucei.
Trypanosoma brucei
Eukaryota; Euzlenzoa; Kinetoplastida; Trypanosomatidae;

REFERENCE
1 (bases 1 to 50)

AUTHORS
Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
Melville, S.E., Rajandream, M.A. and Barrell, B.G.

TITLE
JOURNAL

COMMENT

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridgeshire CB10 1SA, E-mail: barrell@sanger.ac.uk and nh@sanger.ac.uk
Constructed at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (4 kb). The v + i method used for the library construction is described in detail in Smith, H. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaudin and B. Barrell, Oxford University Press, 1999).
Email: nh@sanger.ac.uk
Details of T. brucei sequencing at the Sanger Centre are available at http://www.sanger.ac.uk/projects/T_brucei/.

FEATURES
SOURCE

1..50

```

/organism="Trypanosoma brucei"
/db_xref="taxon:5691"
/clone="154f10"
BASE COUNT      20 a      7 c      9 g      14 t
ORIGIN

Query Match      15.6%; Score 14; DB 13; Length 50;
Best Local Similarity 100.0%; Pred. No. 2.6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atcaaaaaaatatc 14
    |||
Db 10 ATGAAAAAATATAT 23

RESULT 2
AZ829725      25 bp      DNA      GSS      20-FEB-2001
LOCUS
DEFINITION 2M0107124F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0107124 F, DNA sequence.
ACCESSION
VERSION      AZ829725.1 GI:12999549
KEYWORDS
SOURCE      house mouse.
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
1 (bases 1 to 25)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
Plasmid Inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0107 row: I column: 24
Seq primer: CATTGTAAACGACGCCACT
Class: plasmid ends
High quality sequence stop: 25.
Location/Qualifiers
1..25
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0107124"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g114732114|g114f129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated

```

```

with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
BASE COUNT      14 a      3 c      4 g      4 t
ORIGIN

Query Match      14.4%; Score 13; DB 13; Length 25;
Best Local Similarity 100.0%; Pred. No. 9.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 34 ttatgtttcat 46
    |||
Db 24 TTATGTTTTCAT 12

RESULT 3
TA123B12Q/c      26 bp      DNA      GSS      13-DEC-2000
LOCUS
DEFINITION T. brucei sheared genomic DNA clone 123b12, reverse sequence,
genomic survey sequence.
ACCESSION
VERSION      AL463522
KEYWORDS
SOURCE      Trypanosoma brucei.
ORGANISM
Eukaryota; Euzlenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
1 (bases 1 to 26)
Hall,N., Bowman,S., Lennard,N.J., Doggett,J., Atkin,R.,
Chillingworth,C., Ormond,D., Harris,B., El-Sayed,N., Hou,L.,
Melville,S.E., Rajandream,M.A. and Barrell,B.G.
Direct Submission
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
Project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
nhs@sanger.ac.uk
Constructed at the Institute for Genomic Research (TIGR),
Rockville, MD. Genomic DNA isolated from a cloned population of
Trypanosoma brucei (TREU927/4 Gutat 10.1) was mechanically sheared
to give a tight size distribution (
4 kb). The v + i method used for the library construction is
described in detail in Smith, H. and Venter, J.C. (Making small
insert libraries for whole genome shotgun sequencing projects. In
Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
Barrell, Oxford University Press, 1999).
Email: nelsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available
at http://www.sanger.ac.uk/projects/T_brucei/.
Location/Qualifiers
1..26
/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="123b12"
BASE COUNT      12 a      0 c      0 g      14 t
ORIGIN

Query Match      14.4%; Score 13; DB 13; Length 26;
Best Local Similarity 100.0%; Pred. No. 9.5e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaaatattc 16
    |||
Db 20 AAAAAAATATATT 8

RESULT 4
AZ452653      28 bp      DNA      GSS      04-OCT-2000
LOCUS

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DEFINITION 1M0252E07R Mouse 10kb plasmid U06C1M library Mus musculus genomic
 ACCESSION clone U06C1M0252E07 R, DNA sequence.
 VERSION AZ452653
 KEYWORDS AZ452653.1 GI:10609676
 SOURCE GSS.
 ORGANISM house mouse.
 Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murine; Mus.
 REFERENCE 1 (bases 1 to 28)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert length: 10000 Std Error: 0.00
 Plate: 0252 row: E column: 07
 Seq primer: CACACAGGAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 28.
 Location/Qualifiers
 1..28
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="U06C1M0252E07"
 /clone_lib="Mouse 10kb plasmid U06C1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of PMD42 (g11473214(gb)AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."
 BASE COUNT 14 a 2 c 2 g 10 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 13; Length 28;
 Best Local Similarity 100.0%; Pred. No. 9.3e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 4 aaaaaattattt 16
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 Db 16 AAAAAATTATT 28
 RESULT 5
 AZ314238

LOCUS AZ314238 36 bp DNA GSS 29-SEP-2000
 DEFINITION 1M0030N24R Mouse 10kb plasmid U06C1M library Mus musculus genomic
 clone U06C1M0030N24 R, DNA sequence.
 ACCESSION AZ314238
 VERSION AZ314238.1 GI:10359929
 KEYWORDS GSS.
 SOURCE house mouse.
 Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murine; Mus.
 REFERENCE 1 (bases 1 to 36)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert length: 10000 Std Error: 0.00
 Plate: 0030 row: N column: 24
 Seq primer: CACACAGGAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 36.
 Location/Qualifiers
 1..36
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="U06C1M0030N24"
 /clone_lib="Mouse 10kb plasmid U06C1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of PMD42 (g11473214(gb)AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."
 BASE COUNT 8 a 3 c 5 g 20 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 13; Length 36;
 Best Local Similarity 100.0%; Pred. No. 8.5e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 45 atgtttcttatt 57
 |||||||||
 Db 2 ATGTTTCTTATT 14
 RESULT 6
 AZ314238

AA500776 49 bp mRNA EST 01-JUL-1997
 LOCUS AA500776
 DEFINITION vgl01b1.r1 Soares mouse NBHM Mus musculus cDNA clone IMAGE:66061
 5. Similar to TR:G133638 G133638 PARADOXASE 2. ; mRNA sequence.
 ACCESSION AA500776
 VERSION AA500776.1 GI:2235743
 KEYWORDS EST.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 49)
 REFERENCE Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubouque,T.,
 Giesel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
 Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
 Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
 Waterston,R.
 TITLE The WashU-HHMI Mouse EST Project
 JOURNAL Unpublished (1996)
 COMMENT Contact: Marra M/Mouse EST Project
 WashU-HHMI Mouse EST Project
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through LNL; contact the
 IMAGE Consortium (info@image.lnl.gov) for further information.
 MG1:504149
 Trace considered overall poor quality
 Possible reversed clone; similarity on wrong strand
 Seq primer: -28m13 rev2 ET from Amersham
 High quality sequence stop: 1.
 location/Qualifiers
 1..49
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="IMAGE:860061"
 /clone_lib="Soares mouse NBHM"
 /sex="male"
 /tissue_type="heart"
 /dev_stage="4 weeks"
 /lab_host="DH10B"
 /note="Vector: pRT3D-Pac (Pharmacia) with a modified
 polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
 was primed with a Not I - oligo(dT) primer; 15'
 TGTATACCAATCTGAGTGAGGAGCGCCGCAAGATTTTGTGTGTGTGTGT
 3'; double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not
 I and Eco RI sites of the modified pRT3D vector. RNA
 provided by Dr. Minoru KO, Wayne State Univ. Library
 constructed and normalized by Bento Soares and M.Fatima
 Bonaldo."
 BASE COUNT 17 a 8 c 10 g 14 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 10; Length 49;
 Best Local Similarity 100.0%; Pred. No. 7.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 30 aattttatgttt 42
 |||||||
 Db 25 AATTTTATGTTT 37
 RESULT 7
 LOCUS AU102750 50 bp mRNA EST 05-APR-2001
 DEFINITION AU102750 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
 CAS10730, mRNA sequence.
 ACCESSION AU102750

VERSION AU102750.1 GI:13552271
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 1 (bases 1 to 50)
 REFERENCE Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata
 H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo
 K., Suyama,A. and Sugano,S.
 TITLE Fine structural analysis of transcription start sites of human
 mRNAs using full-length enriched and 5'-end enriched cDNA libraries
 JOURNAL Unpublished (2001)
 COMMENT Contact: Yutaka Suzuki
 Department of Virology
 Institute of Medical Science, University of Tokyo
 4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
 Email: yusuzuki@ims.u-tokyo.ac.jp
 Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
 S. Construction and characterization of a full length-enriched and
 a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
 location/Qualifiers
 1..50
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="CAS10730"
 /clone_lib="Sugano Homo sapiens cDNA library"
 BASE COUNT 19 a 13 c 8 g 10 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 10; Length 50;
 Best Local Similarity 100.0%; Pred. No. 7.6e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 73 ccaatgaagcaa 85
 |||||||
 Db 8 CCAATGAAGCAA 20
 RESULT 8
 LOCUS A2579583/C 23 bp DNA GSS 13-DEC-2000
 DEFINITION IM0367N03F Mouse 10kb plasmid UGCGM library Mus musculus genomic
 clone UGCGM0367N03 F, DNA sequence.
 ACCESSION A2579583
 VERSION A2579583.1 GI:11694012
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 23)
 REFERENCE Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0367 row: N column: 03
 Seq primer: CGTGTAAACGACGCGCCAGT
 Class: plasmid ends
 High quality sequence stop: 23.

FEATURES
source

Location/Qualifiers

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1. .23
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C1M0367N03"
/clone_lib="Mouse 10kb plasmid U08C1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (g1147321419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

BASE COUNT
ORIGIN

```
9 a 2 c 0 g 12 t
```

Query Match 13.3%; Score 12; DB 13; Length 23;

Best Local Similarity 100.0%; Pred. No. 2.9e+05;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 aaaaattatt 15

Db 15 AAAAAATTATT 4

RESULT 9
A2996858/c

LOCUS A2996858 24 bp DNA GSS 27-APR-2001

DEFINITION 2M0283L08F Mouse 10kb plasmid U08C2M library Mus musculus genomic

ACCESSION A2996858

VERSION A2996858.1 GI:13868085

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 24)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

TITLE plasmid inserts

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss

University of Utah Genome Center

Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0283 row: L column: 08

Seq primer: CGTGTAAACGACGCCACGT

Class: plasmid ends

FEATURES
sourceHigh quality sequence stop: 24.
Location/Qualifiers

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1. .24
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C2M0283L08"
/clone_lib="Mouse 10kb plasmid U08C2M library"
/sex="Female"
/lab_host="E. coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (female) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (g1147321419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

BASE COUNT
ORIGIN

```
12 a 4 c 3 g 5 t
```

Query Match 13.3%; Score 12; DB 13; Length 24;

Best Local Similarity 100.0%; Pred. No. 2.8e+05;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 33 ttatatgttttc 44

Db 21 TTTATGTTTTC 10

RESULT 10
A2345616/c

LOCUS A2345616 25 bp DNA GSS 29-SEP-2000

DEFINITION 1M0080E20F Mouse 10kb plasmid U08C1M library Mus musculus genomic

ACCESSION A2345616

VERSION A2345616.1 GI:10424853

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 25)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

TITLE plasmid inserts

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss

University of Utah Genome Center

Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0080 row: E column: 20

Seq primer: CGTGTAAACGACGCCACGT

Class: plasmid ends
High quality sequence stop: 25.
Location/Qualifiers

1. 25
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0080E20"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 (g1473214|9b|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
16 a 1 c 0 g 8 t

Query Match 13.3%; Score 12; DB 13; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.8e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 29 taattttttgt 40
|||||
Db 12 TAATTTTATGT 1

RESULT 11
AM246455 27 bp mRNA EST 07-JAN-2000
LOCUS 2821693.3prime NIH_MGC_7 Homo sapiens cDNA clone IMAGE:2821693 3',
DEFINITION mRNA sequence.
ACCESSION AM246455
VERSION AM246455.1 GI:6589448
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 27)
NIH-MGC http://mgc.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished (1999)
Other_ESTS: 2821693.5prime
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue procurement: DCTD/DPF CDNA Library Preparation: Ling Hong/Rubin Laboratory CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNU) DNA Sequencing by: Berkeley MGC sequencing project Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNU at: www.bio.lnu.edu/bhrp/image/image.html Base Calling / Quality Scores: PHRED from University of Washington Genome Center. Vector Trimming: cross_match from University of Washington Genome Center PHRAP suite. Poly-T Identification: patch.pl from Berkeley Drosophila Genome Project. University of Washington Genome Center: http://www.genome.washington.edu Low Quality Sequence: 10

contiguous PHRED high quality bases following vector sequence. Very low quality sequence: Trace file contained 27 contiguous distinct peaks following vector sequence. Polyadenylation: Based upon the presence of a XhoI site followed by a run of 14 or more T residues at the beginning of the sequence, this cDNA insert was polyadenylated.
Plate: LICM7 row: H column: 14
High quality sequence stop: 10.
Location/Qualifiers

1. 27
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2821693"
/clone_lib="NIH MGC 7"
/tissue_type="small cell carcinoma"
/cell_line="MGC3"
/lab_host="DH10B (phage-resistant)"
/note="Organ: lung; Vector: pORF7; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAC(G). Size selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the Laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."

BASE COUNT
ORIGIN
2 a 1 c 2 g 22 t

Query Match 13.3%; Score 12; DB 10; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.7e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 40 ttttcagtgtt 51
|||||
Db 7 TTTTCATGTTT 18

RESULT 12
A2610538 30 bp DNA GSS 13-DEC-2000
LOCUS 1M0435E23R Mouse 10kb plasmid UUCG1M library Mus musculus genomic
DEFINITION clone UUCG1M0435E23 R, DNA sequence.
ACCESSION A2610538
VERSION A2610538.1 GI:11732728
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sclurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 30)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0435 row: E column: 23
Seq primer: CACACAGCAACACCTATGACC
Class: plasmid ends
High quality sequence stop: 30.
Location/Qualifiers
1. 30

```
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0435E23"
/clone_1lb="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-."
/notes="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD42 (g1147321141gb/AP129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

```
source
1. 33
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0349021"
/clone_1lb="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-."
/notes="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD42 (g1147321141gb/AP129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

```
Query Match 13.3%; Score 12; DB 13; Length 30;
Best Local Similarity 100.0%; Pred. No. 2.6e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 27 tgtatttttt 38
|||||
Db 4 TGTATTTTAT 15
```

```
Query Match 13.3%; Score 12; DB 13; Length 33;
Best Local Similarity 100.0%; Pred. No. 2.5e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 40 ttttcatgtttt 51
|||||
Db 16 TTTTCATGTTT 27
```

```
RESULT 13
AZ507698 33 bp DNA GSS 05-OCT-2000
LOCUS
DEFINITION
IM0349021F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0349021 F, DNA sequence.
ACCESSION
AZ507698
VERSION
AZ507698.1 GI:10689014
KEYWORDS
GSS.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
```

```
RESULT 14
AZ487251 38 bp DNA GSS 05-OCT-2000
LOCUS
DEFINITION
IM0316A18R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0316A18 R, DNA sequence.
ACCESSION
AZ487251
VERSION
AZ487251.1 GI:10654814
KEYWORDS
GSS.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
```

```
REFERENCE
1 (bases 1 to 33)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
and Wright,D., Weis,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
```

```
JOURNAL
COMMENT
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
```

```
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Seq Plate: 0349 row: 0 column: 21
Class: plasmid ends
High quality sequence stop: 33.
```

```
Location/Qualifiers
```

FEATURES

```
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Seq Plate: 0316 row: A column: 18
Class: plasmid ends
High quality sequence stop: 38.
```

FEATURES
source

Location/Qualifiers
1. 38
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0316A18"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
14 a 14 c 5 g 5 t

Query Match
Best Local Similarity 100.0%; Score 12; DB 13; Length 38;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 2 tgaataaacta 13
|||||

Db 26 TCAAAAAAATTA 37

RESULT 15
A2662545 41 bp DNA GSS 14-DEC-2000
LOCUS A2662545/c
DEFINITION 1M0541P07R Mouse 10kb plasmid UUCG1M library Mus musculus genomic clone UUCG1M0541P07 R, DNA sequence.
ACCESSION A2662545
VERSION A2662545.1 GI:11799691
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 41)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tinney, A., von Niederhausern, A. and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunne@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0541 row: P column: 07
Seq primer: CACACAGGAACACGCTATGACC
Class: plasmid ends

FEATURES
source

High quality sequence stop: 41.
Location/Qualifiers
1. 41
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0541P07"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
16 a 9 c 5 g 11 t

Query Match
Best Local Similarity 100.0%; Score 12; DB 13; Length 41;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 52 cttatgttga 63
|||||

Db 35 CTTATTGTTGA 24

RESULT 16
T17635 42 bp mRNA EST 06-JUN-1994
LOCUS T17635/c
DEFINITION mps v360 The blue guys library Saccharomyces cerevisiae cDNA sequence upstream of lacZ fusion similar to GAC1, X63941, mRNA sequence.
ACCESSION T17635
VERSION T17635.1 GI:459560
KEYWORDS EST.
SOURCE baker's yeast.
ORGANISM Saccharomyces cerevisiae
Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes; Saccharomycetales; Saccharomycetaceae; Saccharomyces. 1 (bases 1 to 42)
Burns, N., Grimwade, B., Ross-Macdonald, P. B., Choi, E.-Y., Finberg, K., Roeder, G. S. and Snyder, M.
Large-scale analysis of gene expression, protein localization and gene disruption in Saccharomyces cerevisiae
Genes Dev. 8, 1087-1105 (1994)
95011603
Contact: Snyder M
Department of Biology
Yale University
New Haven CT 06520-8103
Tel: 2034326139
Fax: 2034326161
Email: snymp@yalevm.ycc.yale.edu
lacZ fusion; Vegetative expression; Beta-gal fusion localization pattern;
50 cytoplasmic spots; Disruption phenotype: none detected; Fusion: codon 407 of GAC1 gene. Sequence below near or adjacent to lacZ.

Seq primer: lacZ sequences in transposon.
Location/Qualifiers

1. 42
/organism="Saccharomyces cerevisiae"
/db_xref="taxon:4932"
/clone_id="The blue guys library"
/lab_host="E.coli"

/note="Vector: PRECmtn: A yeast genomic DNA library was prepared in the vector pHS6, and subjected to transposon mutagenesis with mtn3. This mini-transposon carries lacZ sequences that lack an initiation codon; expression of lacZ is only provided by in frame fusion to yeast coding sequence. The yeast genomic DNA carrying the transposon was excised from pHS6 and transplanted back onto the yeast chromosome. Yeast colonies expressing lacZ were screened for in a color assay. A plasmid containing the genomic DNA/lacZ fusion junction was recovered from each individual yeast colony that expressed lacZ activity. These recovered plasmids comprise 'The blue guys library'. The fusion junction was then sequenced to identify the expressed ORF upstream of the fusion."

BASE COUNT
ORIGIN
7 a 12 c 11 g 12 t

Query Match 13.3%; Score 12; DB 11; Length 42;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 75 aatgaagcaag 86
|||||
Db 22 AATGAAGCAAG 11

RESULT 17
AZ514394

LOCUS 45 bp DNA GSS 05-OCT-2000
DEFINITION 1M0361J02F Mouse 10kb plasmid UNGC1M library Mus musculus genomic
clone UNGC1M0361J02 F, DNA sequence.

ACCESSION AZ514394
VERSION AZ514394.1 GI:10695710
KEYWORDS GSS.

SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciuromorphi; Muridae; Murinae; Mus.
1 (bases 1 to 45)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmood,M., Meenen,E., Pedersen,T., Reilly,
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA

Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0361 row: 5 column: 02
Seq primer: CGTGTAAACGACGCCACG
Class: plasmid ends
High quality sequence stop: 45.
Location/Qualifiers

1. 45
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UNG1M0361J02"

/clone_id="Mouse 10kb plasmid UNGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PMD42ny: Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g1147321141gb1AF129072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 24 a 2 c 9 g 10 t
ORIGIN

Query Match 13.3%; Score 12; DB 13; Length 45;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaaatatt 15
|||||
Db 3 AAAAAATTATT 14

RESULT 18
A1805932

LOCUS 46 bp mRNA EST 13-DEC-1999
DEFINITION t622903.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
IMAGE:2090356 3 similar to TR:Q99766 Q99766 HYPOTHETICAL 15.7 KD
PROTEIN: ; mRNA sequence.

ACCESSION A1805932
VERSION A1805932.1 GI:5392498
KEYWORDS EST.

SOURCE human.
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 46)
NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

TITLE Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov

This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 606 Std Error: 0.00
Seq primer: -40UP from Gldco
High quality sequence stop: 1.
Location/Qualifiers

1. 46
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2090356"
/clone_id="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT73D-Pac (Pharmacia) with
a modified polylinker; Site:1: Not I; Site:2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NBH19W, testis NHT, and B-cell
NCI-CCAP_GCB1) were mixed, and ss circles were made in

vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 297480-302087, 682632-687239, 726408-728711, and 729096-731399. Subtraction by Bento Soares and M. Fatima Bonaldo.

BASE COUNT 13 a 9 c 5 g 19 t
ORIGIN

Query Match 13.3%; Score 12; DB 10; Length 46;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 25 actgaattttt 36
Db 34 ACTGTAATTTT 45

RESULT 19
AZ832106/c

LOCUS 48 bp DNA GSS 20-FEB-2001
DEFINITION 2M0112113F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0112113 F, DNA sequence.

ACCESSION AZ832106
VERSION AZ832106.1 GI:13002014

KEYWORDS GSS.

SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 48)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLCT, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177

Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0112 row: 1 column: 13
Seq primer: CGTTGTAACGACGCGCAGT

Class: plasmid ends
High quality sequence stop: 48.

FEATURES
SOURCE location/Qualifiers
1. 48

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0112113"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative of PMD42 (GI14732114|gb|AF129072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 14 a 11 c 3 g 20 t
ORIGIN

Query Match 13.3%; Score 12; DB 13; Length 48;
Best Local Similarity 100.0%; Pred. No. 2.2e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaattatt 15
Db 48 AAAAAATTATT 37

RESULT 20
D19972/c

LOCUS 50 bp mRNA EST 30-JUL-1996
DEFINITION HUMGS00937 Human promyelocyte Homo sapiens cDNA clone mm06d08 3',
mRNA sequence.

ACCESSION D19972
VERSION D19972.1 GI:500869

KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo. 1 (bases 1 to 50)

AUTHORS Okubo,K., Fukushima,A., Yoshii,J., Niyama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
Gene expression of human promyelocytic cell line HL60 before and after induction of differentiation. A new application of 3'directed cDNA sequencing

TITLE Unpublished (1993)

JOURNAL

COMMENT Contact: Okubo,K., Fukushima,A., Yoshii,J., Niyama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
Institute for Molecular and Cellular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.

FEATURES
SOURCE location/Qualifiers
1. 50

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="mm06d08"
/clone_lib="Human promyelocyte"
/note="Female, adult, cell_line = HL60, cell_type = promyelocyte."

BASE COUNT 20 a 5 c 11 g 14 t
ORIGIN

Query Match 13.3%; Score 12; DB 11; Length 50;
Best Local Similarity 100.0%; Pred. No. 2.2e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 ttcaaatgttac 26
Db 20 TTCAAAAGTTTAC 9

RESULT 21
AZ782616

LOCUS 20 bp DNA GSS 16-FEB-2001
DEFINITION 2M0023F17R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0023F17 R, DNA sequence.

ACCESSION AZ782616
VERSION AZ782616.1 GI:12916517

KEYWORDS
GSS.
house mouse.
SOURCE
Mus musculus
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 20)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
AUTHORS
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
TITLE
Unpublished (2000)
JOURNAL
Contact: Robert B. Weiss
University of Utah
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0023 row: F column: 17
Seq primer: CACACAGCAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 20.
FEATURES
Location/Qualifiers
1..20
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG2M0023F17"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gb]AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
BASE COUNT
6 a 4 c 1 g 9 t
ORIGIN
Query Match 12.2%; Score 11; DB 13; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.8e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 33 ttttatgttt 43
|||||
Db 9 ttttatgttt 19
RESULT 22
A2768088 21 bp DNA GSS 16-FEB-2001
LOCUS A2768088
DEFINITION IM0567G21R Mouse 10kb plasmid UUCG1M library Mus musculus genomic
clone UUCG1M0567G21 R, DNA sequence.
ACCESSION A2768088

VERSION
A2768088.1 GI:12886839
KEYWORDS
GSS.
house mouse.
SOURCE
Mus musculus
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 21)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
AUTHORS
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
TITLE
Unpublished (2000)
JOURNAL
Contact: Robert B. Weiss
University of Utah
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0567 row: G column: 21
Seq primer: CACACAGCAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 21.
FEATURES
Location/Qualifiers
1..21
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0567G21"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gb]AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
BASE COUNT
5 a 3 c 1 g 12 t
ORIGIN
Query Match 12.2%; Score 11; DB 13; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.8e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 40 tttcatgttt 50
|||||
Db 11 tttcatgttt 21
RESULT 23
AA991150 22 bp mRNA EST 03-JUN-1998
LOCUS AA991150/c
DEFINITION os40a07.s1 NCI CGAP Br2 Homo sapiens cDNA clone IMAGE:1607796 3'
similar to TR:Q35990 Q35990 HYPOTHETICAL 8.9 KD PROTEIN.; contains

element L1 repetitive element ;, mRNA sequence.

ACCESSION AA991150
 VERSION AA991150.1 GI:3177639
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 1 (bases 1 to 22)
 NCI-CCAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 Unpublished (1997)
 JOURNAL Contact: Robert Strausberg, Ph.D.
 COMMENT Email: cgapsb@mail.nih.gov
 Tissue Procurement: Christopher Moskalko, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CCAP clone distribution information can be found through the I.M.A.G.E. Consortium/ULNL at: www.bio.lnlnl.gov/bdip/image/image.html

Trace considered overall poor quality
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 1.
 Location/Qualifiers
 1..22
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:1607796"
 /clone_lib="NCI-CCAP_Br2"
 /sex="female, pooled"
 /tissue_type="breast"
 /lab_host="DH10B"
 /note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from pooled bulk breast tumor tissue, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. This library is the normalized version of NCI-CCAP_Br1.1. Library was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 16 a 3 c 2 g 1 t
 ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 22;
 Best Local Similarity 100.0%; Pred. No. 8.5e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 46 tgtttcttat 56
 |||||||||
 Db 22 TGTTCCTTAT 12

RESULT 24
 A2766483 22 bp DNA GSS 16-FEB-2001
 LOCUS A2766483
 DEFINITION 1M056410F Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M056410 F, DNA sequence.
 ACCESSION A2766483
 VERSION A2766483.1 GI:12883604
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sclerothelmi; Muridae; Murinae; Mus.
 1 (bases 1 to 22)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly

, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A. and Wright, D., Weiss, R.
 Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 Unpublished (2000)
 JOURNAL Contact: Robert B. Weiss
 COMMENT University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunne@genetics.utah.edu
 Insert Length: 1000 Std Error: 0.00
 Plate: 0564 row: A column: 10
 Seg primer: CGTGTAAACGACGCCACGT
 Class: plasmid ends
 High quality sequence stop: 22.
 Location/Qualifiers
 1..22
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M056410"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: pMD42uv: Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g1473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 4 a 2 c 4 g 12 t
 ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 22;
 Best Local Similarity 100.0%; Pred. No. 8.5e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 46 tgtttcttat 56
 |||||||||
 Db 4 TGTTCCTTAT 14

RESULT 25
 A2387817 23 bp DNA GSS 02-OCT-2000
 LOCUS A2387817
 DEFINITION 1M0147824R Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0147824 R, DNA sequence.
 ACCESSION A2387817
 VERSION A2387817.1 GI:10501525
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sclerothelmi; Muridae; Murinae; Mus.
 1 (bases 1 to 23)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,

Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A., and Wright, D., Weiss, R.
 Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL
 COMMENT
 Unpublished (2000)
 Contact: Robert B. Weiss
 University of Utah Genome Center
 Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunne@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0147 row: B column: 24
 Seq primer: CACACAGCAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 23.

FEATURES
 SOURCE
 Location/Qualifiers

1. .23
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="U08C1M0147B24"
 /clone_lib="Mouse 10kb plasmid U08C1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g1147321149b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
 ORIGIN
 5 a 4 c 3 g 11 t
 Query Match 12.2%; Score 11; DB 13; Length 23;
 Best Local Similarity 100.0%; Pred. No. 8.4e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 28 gtaattttat 38
 ||||||||||||
 Db 6 gtaattttat 16

RESULT 26
 TA242F03P 24 bp DNA GSS 13-DEC-2000
 LOCUS
 DEFINITION T. Brucei sheared genomic DNA clone 242F03, forward sequence, genomic survey sequence.
 ACCESSION
 VERSION AL482984
 KEYWORDS AL482984.1 GI:11848725
 SOURCE GSS.
 ORGANISM Trypanosoma brucei.
 Trypanosoma brucei
 Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
 Trypanosoma.
 REFERENCE 1 (bases 1 to 24)
 Location/Qualifiers

AUTHORS
 TITLE
 JOURNAL
 COMMENT
 Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R., Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L., Melville, S.E., Rajandream, M.A. and Barrall, B.G.
 Direct Submission
 Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, E-mail: barrall@sanger.ac.uk and nh@sanger.ac.uk

Constructed at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 G07at 10.1) was mechanically sheared to give a tight size distribution (4 kb). The v + i method used for the library construction is described in detail in Smith, H. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaubin and B. Barrall, Oxford University Press, 1999).
 Email: neilsayed@tigr.org
 Details of T. brucei sequencing at the Sanger Centre are available at <http://www.sanger.ac.uk/projects/T-brucei/>.

FEATURES
 SOURCE
 Location/Qualifiers

1. .24
 /organism="Trypanosoma brucei"
 /strain="TREU927"
 /db_xref="taxon:5691"
 /clone="242f03"
 BASE COUNT
 ORIGIN
 1 a 3 c 10 g 10 t
 Query Match 12.2%; Score 11; DB 13; Length 24;
 Best Local Similarity 100.0%; Pred. No. 8.2e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 52 ctattgtttg 62
 ||||||||||||
 Db 8 cttattgtttg 18

RESULT 27
 LOCUS
 DEFINITION A2462654 25 bp DNA GSS 04-OCT-2000
 A2462654
 clone U08C1M0269P09 R, DNA sequence.
 ACCESSION
 VERSION A2462654.1 GI:10620695
 KEYWORDS GSS.
 SOURCE
 ORGANISM Mus musculus.
 house mouse.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 I (bases 1 to 25)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A., and Wright, D., Weiss, R.
 Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 Unpublished (2000)
 Contact: Robert B. Weiss
 University of Utah Genome Center
 Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunne@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0269 row: P column: 09
 Seq primer: CACACAGCAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 25.
 Location/Qualifiers

FEATURES
 SOURCE
 Location/Qualifiers

source

1. .25
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C2M0073120"
/clone_lib="Mouse 10Kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv: Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
10 a 2 c 0 g 13 t

Query Match
Best Local Similarity 12.2%; Score 11; DB 13; Length 25;
Matches 11: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 6 aaaaattattt 16
|||||
Db 4 AAAAATTATT 14

RESULT 28
A2809415 28 bp DNA GSS 20-FEB-2001
LOCUS A2809415
DEFINITION 2M0073120F Mouse 10kb plasmid U06C1M library Mus musculus genomic clone U06C2M0073120 F, DNA sequence.
ACCESSION A2809415
VERSION A2809415.1 GI:12975693
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sclerothnathi; Muridae; Murinae; Mus.
1 (bases 1 to 28)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmood, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A., and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10Kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0073 row: 1 column: 20
Seq primer: CGTTTAAACGACGCGCAGT
Class: plasmid ends
High quality sequence stop: 28.

FEATURES
source

Location/Qualifiers
1. .28
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C2M0073120"
/clone_lib="Mouse 10Kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv: Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
7 a 5 c 5 g 11 t

Query Match
Best Local Similarity 12.2%; Score 11; DB 13; Length 28;
Matches 11: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 74 caatgaagca 84
|||||
Db 14 CAATGAAGCA 4

RESULT 29
D18726 29 bp mRNA EST 12-DEC-1995
LOCUS D18726
DEFINITION MUSGS01788 Mouse 3'-directed Mus musculus domesticus cDNA clone md0169 3', mRNA sequence.
ACCESSION D18726
VERSION D18726.1 GI:1100695
KEYWORDS EST.
SOURCE Western European house mouse.
ORGANISM Mus musculus domesticus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sclerothnathi; Muridae; Murinae; Mus.
1 (bases 1 to 29)
Kawamoto, S., Okubo, K., Yoshii, J., Katsuki, M., and Matsubara, K.
Analysis of gene expression in mouse embryogenesis by 3'-directed cDNA sequencing
JOURNAL Unpublished (1995)
COMMENT Contact: Kawamoto, S., Okubo, K., Yoshii, J., Katsuki, M. and Matsubara, K.
Institute for Cellular and Molecular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.
FEATURES
source
1. .29
/organism="Mus musculus domesticus"
/strain="C57BL/6J"
/db_xref="taxon:10092"
/clone="md0169"
/clone_lib="Mouse 3'-directed"
/tissue_type="decidual tissue (day 6.5-8.5 of gestation)"
BASE COUNT
ORIGIN
15 a 3 c 2 g 9 t

Query Match 12.2%; Score 11; DB 11; Length 29;
 Best Local Similarity 100.0%; Pred. No. 7.7e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 36 tatgttttcac 46
 |||||||
 Db 15 TATGTTTCAT 25

RESULT 30

AZ580321 29 bp DNA GSS 13-DEC-2000
 LOCUS AZ580321
 DEFINITION IM0366G02R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0366G02 R, DNA sequence.
 ACCESSION AZ580321
 VERSION AZ580321.1 GI:11694750
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 29)
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 'M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

TITLE Unpublished (2000)
 JOURNAL Contact: Robert B. Weiss
 COMMENT University of Utah
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunne@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0368 row: G column: 02
 Seq primer: CACACAGAAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 29.

FEATURES
 SOURCE Location/Qualifiers
 1..29
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0366G02"
 /clone_1lb="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pMD2 (gii14732114[gb]IAFI29072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

BASE COUNT 10 a 11 c 3 g 5 t

ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 29;
 Best Local Similarity 100.0%; Pred. No. 7.7e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 76 atgaagcaag 86
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 Db 16 ATGAAGCAAG 26

RESULT 31

AZ579378 31 bp DNA GSS 13-DEC-2000
 LOCUS AZ579378
 DEFINITION IM0363N13R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0363N13 R, DNA sequence.
 ACCESSION AZ579378
 VERSION AZ579378.1 GI:11693807
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 31)
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 'M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

TITLE Unpublished (2000)
 JOURNAL Contact: Robert B. Weiss
 COMMENT University of Utah
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunne@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0363 row: N column: 13
 Seq primer: CACACAGAAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 31.

FEATURES
 SOURCE Location/Qualifiers
 1..31
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0363N13"
 /clone_1lb="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pMD2 (gii14732114[gb]IAFI29072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

```

BASE COUNT      12 a      1 c      8 g      10 t
ORIGIN
Query Match      12.2%; Score 11; DB 13; Length 31;
Best Local Similarity 100.0%; Pred. No. 7.5e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY      4 aaaaaaatat 14
Db      3 AAAAAATTTAT 13

RESULT 32
AZ331642/c      32 bp      DNA      GSS      29-SEP-2000
LOCUS
DEFINITION      1M0059P1R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
ACCESSION      AZ331642
VERSION      AZ331642.1 GI:10394528
KEYWORDS
SOURCE      house mouse.
ORGANISM      Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 32)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0059 row: P column: 11
Seq primer: CACACGAGAAACAGCTAGGACC
Class: plasmid ends
High quality sequence stop: 32.
Location/Qualifiers
1..32
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0059P1"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, T1-resistant, F-"
/notes="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (911473211419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells

```

```

BASE COUNT      14 a      7 c      1 g      10 t
ORIGIN
Query Match      12.2%; Score 11; DB 13; Length 32;
Best Local Similarity 100.0%; Pred. No. 7.5e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY      13 attccaagt 23
Db      20 ATTTCAAAGTT 10

RESULT 33
AZ591923      32 bp      DNA      GSS      13-DEC-2000
LOCUS
DEFINITION      1M0402F21F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
ACCESSION      AZ591923
VERSION      AZ591923.1 GI:11714113
KEYWORDS
SOURCE      house mouse.
ORGANISM      Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 32)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0402 row: F column: 21
Seq primer: GGTGTAAACGACGCCAGCT
Class: plasmid ends
High quality sequence stop: 32.
Location/Qualifiers
1..32
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0402F21"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, T1-resistant, F-"
/notes="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (911473211419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into

```


chemically-competent *E. coli* XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 8 a 5 c 9 g 10 t
ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 32;
Best Local Similarity 100.0%; Pred. No. 7.5e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 34
LOCUS U44209/c 33 bp mRNA EST 03-APR-1996
DEFINITION ENU44209 *Aspergillus nidulans* cleistothecium *Emericella nidulans*
cDNA clone SE0393, mRNA sequence.
ACCESSION U44209
VERSION U44209.1 GI:1244872
KEYWORDS EST.
SOURCE *Emericella nidulans*.
ORGANISM *Emericella nidulans*.
Eukaryota; Fungi; Ascomycota; Pezizomycotina; Eurotiomycetes;
Eurotiiales; Trichocomaceae; *Emericella*.

REFERENCE 1 (bases 1 to 33)
AUTHORS Lee,D., Lee,S., Hwang,H., Kim,J. and Chae,K.
TITLE Quantitative analysis of gene expression in sexual structures of *Aspergillus nidulans* by sequencing of 3'-directed cDNA clones
JOURNAL FEMS Microbiol. Lett. 138 (1), 71-76 (1996)
MEDLINE 96236220
COMMENT Contact: Keon-Sang Chae
Chonbuk National University
Chonju, 561-756, S. Korea
Tel: +82-652-70-3340
Fax: +82-652-70-3345
Email: chaek@chonbukns.chonbuk.ac.kr.
Location/Qualifiers

FEATURES
SOURCE 1. .33
/organism="Emicella nidulans"
/strain="FGSC4"
/db_xref="taxon:162425"
/clone="SE0393"
/clone_lib="Aspergillus nidulans cleistothecium"
/tissue_type="cleistothecium"
/cell_type="Hull cell"
/dev_stage="sexual"
/note="3'-directed cDNA clones; single-pass sequencing"

BASE COUNT 18 a 3 c 2 g 10 t
ORIGIN

Query Match 12.2%; Score 11; DB 11; Length 33;
Best Local Similarity 100.0%; Pred. No. 7.4e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 32 ttattatgtt 42
|||||
Db 31 TTTTATGTTT 21

RESULT 35
LOCUS A2423204 33 bp DNA GSS 03-OCT-2000
DEFINITION c106202010F Mouse 10kb plasmid UNGC1M library Mus musculus genomic
clone UNGC1M0202010 F, DNA sequence.
ACCESSION A2423204
VERSION A2423204.1 GI:10547217
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 33)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0202 row: 0 column: 10
Seq primer: CGTTGTAAACGACGCCACGT
Class: plasmid ends
High quality sequence stop: 33.
Location/Qualifiers

FEATURES
SOURCE 1. .33
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UNG1M0202010"
/clone_lib="Mouse 10kb plasmid UNGC1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42mv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (g114732114[gb|AF129072.1]), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent *E. coli* XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 15 a 3 c 7 g 8 t
ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 33;
Best Local Similarity 100.0%; Pred. No. 7.4e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 aaaaaattat 14
|||||
Db 21 AAAAAATTAT 31

RESULT 36
LOCUS A2494328 33 bp DNA GSS 05-OCT-2000
DEFINITION 1M0329F05R Mouse 10kb plasmid UNGC1M library Mus musculus genomic
clone UNGC1M0329F05 R, DNA sequence.
ACCESSION A2494328
VERSION A2494328.1 GI:10668799
KEYWORDS GSS.
SOURCE house mouse.

ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 33)
 AUTHORS Dun, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tinney, A., von Niederhausen, A., and Wright, D., Weiss, R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT 84112 USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert length: 10000 Std Error: 0.00
 Plate: 0329 row: F column: 05
 Seq primer: CACACAGGAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 33.
 Location/Qualifiers
 1..33
 /organism="Mus musculus"
 /strain="C57Bl/6J"
 /db_xref="taxon:10090"
 /clone="U08C1M0329F05"
 /clone_lib="Mouse 10kb plasmid U08C1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57Bl/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g14732114(gblAF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 8 a 1 c 7 g 17 t
 ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 33;
 Best Local Similarity 100.0%; Pred. No. 7.4e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 49 ttctctatgt 59
 |||||
 Db 23 ttctctatgt 33

RESULT 37
 AU014463 35 bp mRNA EST 03-AUG-1998
 LOCUS AU014463 Schizosaccharomyces pombe late log phase cDNA
 DEFINITION AU014463 Schizosaccharomyces pombe cDNA clone spc09897, mRNA sequence.
 ACCESSION AU014463
 VERSION AU014463.1 GI:3369254
 KEYWORDS EST.

SOURCE fission yeast.
 ORGANISM Schizosaccharomyces pombe
 Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes; Schizosaccharomycetales; Schizosaccharomycetaceae; Schizosaccharomyces.
 REFERENCE 1 (bases 1 to 35)
 AUTHORS Moriyama, M. and Mita, K.
 TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe
 JOURNAL Unpublished (1998)
 COMMENT Contact: Mitsuoki Moriyama
 Genome Research Group
 National Institute of Radiological Sciences
 9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
 Email: moriyoma@nirs.go.jp
 Location/Qualifiers
 1..35
 /organism="Schizosaccharomyces pombe"
 /strain="972"
 /db_xref="taxon:4896"
 /clone="spc09897"
 /clone_lib="Schizosaccharomyces pombe late log phase cDNA"
 /sex="h minus"
 /note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)"

BASE COUNT 21 a 1 c 5 g 8 t
 ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 35;
 Best Local Similarity 100.0%; Pred. No. 7.2e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 gaaaaaatta 13
 |||||
 Db 16 gaaaaaatta 26

RESULT 38
 D19995 35 bp mRNA EST 30-JUL-1996
 LOCUS D19995 Human promyelocyte Homo sapiens cDNA clone mp1322 3', mRNA sequence.
 DEFINITION H06GS00963 Human promyelocyte Homo sapiens cDNA clone mp1322 3', mRNA sequence.
 ACCESSION D19995
 VERSION D19995.1 GI:500892
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 35)
 AUTHORS Okubo, K., Fukushima, A., Yoshii, J., Niijima, T., Kojima, Y., Yoshinari, H., Arimoto, J. and Matsubara, K.
 TITLE Gene expression of human promyelocytic cell line HL60 before and after induction of differentiation. A new application of 3'directed cDNA sequencing
 JOURNAL Unpublished (1993)
 COMMENT Contact: Okubo, K., Fukushima, A., Yoshii, J., Niijima, T., Kojima, Y., Yoshinari, H., Arimoto, J. and Matsubara, K.
 Institute for Molecular and Cellular Biology
 Osaka University
 3-1 Yamada-oka, Suita, Osaka 565, Japan.
 Location/Qualifiers
 1..35
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="mp1322"
 /note="Female, adult, cell_line = HL60, cell_type =

BASE COUNT 14 a 7 c 2 g 12 t
ORIGIN

Query Match 12.2%; Score 11; DB 11; Length 35;
Best Local Similarity 100.0%; Pred. No. 7.2e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 27 tgaattttta 37
|||||
Db 20 TGTATTTTVA 10

RESULT 39
AU111149/c 36 bp mRNA EST 19-OCT-2000
LOCUS AU111149 unpublished oligo-capped cDNA library Caenorhabditis
DEFINITION elegans cDNA yk724e1 5', mRNA sequence.
ACCESSION AU111149
VERSION AU111149.1 GI:10924716
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea
; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 36)
Kohara, Y., Shin-i, T., Thierry-Mieg, J., Thierry-Mieg, D., Suzuki, Y.
and Sugano, S.
A complementary view of the C. elegans genome
Unpublished (2000)
Contact: Yuji Kohara
Genome Biology Lab.
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.
Location/Qualifiers

REFERENCE 1 (bases 1 to 36)
AUTHORS Kohara, Y., Shin-i, T., Thierry-Mieg, J., Thierry-Mieg, D., Suzuki, Y.
and Sugano, S.
A complementary view of the C. elegans genome
Unpublished (2000)
Contact: Yuji Kohara
Genome Biology Lab.
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.
Location/Qualifiers

FEATURES
SOURCE
1. .36
/organism="Caenorhabditis elegans"
/strain="N2"
/db_xref="taxon:6239"
/clone="YK724e1"
/clone_1bp="unpublished oligo-capped cDNA library"
/sex="Hermaphrodite"
/tissue_type="whole animal"
/dev_stage="varied"
BASE COUNT 17 a 3 c 6 g 10 t
ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 36;
Best Local Similarity 100.0%; Pred. No. 7.2e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 aaaaattattt 16
|||||
Db 23 AAAAATTATTT 13

RESULT 40
AZ381596/c 36 bp DNA GSS 02-OCT-2000
LOCUS AZ381596
DEFINITION IM0138C1F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0138C16 F, DNA sequence.
ACCESSION AZ381596
VERSION AZ381596.1 GI:10495296
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus.
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 36)
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0138 row: C column: 16
Seq primer: CGTGTAAACGACGCGCAGT
Class: plasmid ends
High quality sequence stop: 36.
Location/Qualifiers

TITLE
JOURNAL
COMMENT

FEATURES
SOURCE
1. .36
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0138C16"
/clone_1bp="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g114732114[9b]AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 19 a 2 c 10 g 5 t
ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 36;
Best Local Similarity 100.0%; Pred. No. 7.2e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 44 catgttttctt 54
|||||
Db 36 CATGTTTCTT 26

RESULT 41
AZ807406/c 36 bp DNA GSS 20-FEB-2001
LOCUS AZ807406
DEFINITION 2M0070D15F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0070D15 F, DNA sequence.
ACCESSION AZ807406
VERSION AZ807406.1 GI:12971722
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE 1 (bases 1 to 36)
 AUTHORS Dunp,D., Aoyagi,A., Barber,M., Beacorn,T., Duvall,B., Hamill,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausen,A., and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10Kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0070 row: D column: 15
 Seq primer: CGTGTAAACGACGCGCAGT
 Class: plasmid ends
 High quality sequence stop: 36.
 Location/Qualifiers
 1..36
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUCG2M0070D15"
 /clone_lib="Mouse 10Kb plasmid UUCG1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 18 a 7 c 7 g 4 t
 ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 36;
 Best Local Similarity 100.0%; Pred. No. 7.2e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtattttt 36
 |||||||
 Db 19 CTGTAATTTT 9

RESULT 42
 AA647854 37 bp mRNA EST 28-OCT-1997
 LOCUS AA647854
 DEFINITION v980e05.s1 Knowles Solter mouse 2 cell Mus musculus cDNA clone IMAGE:1108640 5' similar to TR:G1136390 G1136390 KIA0164 PROTEIN.
 ; mRNA sequence.
 ACCESSION AA647854
 VERSION AA647854.1 GI:2574283
 KEYWORDS EST.
 SOURCE house mouse.

ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 37)
 AUTHORS Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T., Gesel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M., Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B., Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and Waterston,R.
 TITLE The WashU-HHMI Mouse EST Project
 JOURNAL Unpublished (1996)
 COMMENT Contact: Marra M/Mouse EST Project
 WashU-HHMI Mouse EST Project
 Washington University School of Medicine
 444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through INM; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
 MGI:606808
 Trace considered overall poor quality
 Possible reversed clone; similarity on wrong strand
 High quality sequence stop: 1.
 Location/Qualifiers
 1..37
 /organism="Mus musculus"
 /strain="B6D2 F1/3"
 /db_xref="taxon:10090"
 /clone="IMAGE:1108640"
 /clone_lib="Knowles Solter mouse 2 cell"
 /tissue_type="embryo"
 /dev_stage="2-cell"
 /lab_host="DH10B"
 /note="Organ: embryo; Vector: pBluescribe (modified); Site_1: MluI; Site_2: SalI; Cloned unidirectionally from mRNA prepared from 13,500 2-cell stage embryos. Primer: SalI(dT): 5'-CGGCGACGCGACGCTTTT-3'. cDNAs were cloned into the MluI/SalI sites of a modified pBluescribe vector using commercial linkers (NEB). Average insert size: 1.2 kb."

BASE COUNT 9 a 8 c 8 g 12 t
 ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 37;
 Best Local Similarity 100.0%; Pred. No. 7.1e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 66 aataagtcac 76
 |||||||
 Db 26 AATAAGTCCAA 36

RESULT 43
 AI200438 37 bp mRNA EST 29-NOV-1998
 LOCUS AI200438
 DEFINITION qf93b01.x1 Soares-placenta_8to9weeks_2NBHP6to9M Homo sapiens cDNA clone IMAGE:1757545 3' similar to SW:Q0PT_HUMAN Q16769 GLUTAMINYL-PEPTIDE CYCLOTRANSFERASE PRECURSOR; contains element MER35 repetitive element;; mRNA sequence.
 ACCESSION AI200438
 VERSION AI200438.1 GI:3753044
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 37)
 AUTHORS NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 JOURNAL Unpublished (1997)

COMMENT Contact: Robert Strausberg, Ph.D.
Email: ccaps-remail.nih.gov
This clone is available royalty-free through LNL; contact the IMAGE Consortium (infoimage.lnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 863 Std Error: 0.00
Seq primer: -40UP from GIBCO
High quality sequence stop: 1.
Location/Qualifiers

FEATURES
source
1. .37
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1757545"
/clone_lib="Soares, Placenta, 8to9weeks_2nbHpt09W"
/dex_stage="two placentae: one from 8 weeks and another from 9 weeks post conception"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: placenta; Vector: pT73D (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer (5' TGTTACCAATCTGAGTGGAGGCGCGCGATTTTCTTTTCTTTT 3'), double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library constructed by Bento Soares and M. Fatima Donaldso."

BASE COUNT 11 a 6 c 12 g 8 t

ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 37;
Best Local Similarity 100.0%; Pred. No. 7.1e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 tgaaaaaatt 12
|||||
Db 14 TGAATAAATT 24

RESULT 44
A2806836 37 bp DNA GSS 20-FEB-2001
LOCUS 2M0069108F Mouse 10kb plasmid UUC1M library Mus musculus genomic
DEFINITION clone UUC2M0069108 F, DNA sequence.
ACCESSION A2806836
VERSION A2806836.1 GI:12970584
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murine; Mus.
1 (bases 1 to 37)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0069 row: I column: 08
Seq primer: CGTGTAAACGACGCGCAGC
Class: plasmid ends
High quality sequence stop: 37.
Location/Qualifiers

FEATURES

source

1. .37
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUC2M0069108"
/clone_lib="Mouse 10kb plasmid UUC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (gii4732114(gb)AFL29072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 14 a 1 c 10 g 12 t

ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 37;
Best Local Similarity 100.0%; Pred. No. 7.1e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 30 aattttatgt 40
|||||
Db 5 AATTTTTATGT 15

RESULT 45
D21038 38 bp mRNA EST 30-JUL-1996
LOCUS HMG502021 Human promyelocyte Homo sapiens cDNA clone mp0144 3',
DEFINITION mRNA sequence.
ACCESSION D21038
VERSION D21038.1 GI:504858
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 38)
Okubo,K., Fukushima,A., Yoshii,J., Niijama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
Gene expression of human promyelocytic cell line HL60 before and after induction of differentiation. A new application of 3'directed cDNA sequencing
Unpublished (1993)
Contact: Okubo,K., Fukushima,A., Yoshii,J., Niijama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
Institute for Molecular and Cellular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.
Location/Qualifiers

FEATURES
source
1. .38
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="mp0144"
/clone_lib="human promyelocyte"
/note="Female, adult, cell_line = HL60, cell_type = promyelocyte."
Location/Qualifiers

BASE COUNT 15 a 4 c 8 g 11 t

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 03:22:36 ; Search time 1494.92 Seconds
(without alignments)
993.194 Million cell updates/sec

Title: US-09-531-438-4
Perfect score: 90
Sequence: 1 atgaaaaaattatttcaaa.....gtccaatgaagaagtga 90

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 1472140 seqs, 8248589755 residues

Word size : 0

Total number of hits satisfying chosen parameters: 541028

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: listing first 45 summaries

Database : GenEmbl.*
1: gb_ba.*
2: gb_hvg.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vl.*
15: em_ba.*
16: em_fun.*
17: em_hum.*
18: em_in.*
19: em_om.*
20: em_or.*
21: em_ov.*
22: em_pat.*
23: em_ph.*
24: em_pl.*
25: em_ro.*
26: em_sts.*
27: em_sy.*
28: em_un.*
29: em_vl.*
30: em_hugo_hum.*
31: em_hugo_inv.*
32: em_hugo_rod.*
33: em_hvg_hum.*
34: em_hvg_inv.*
35: em_hvg_rod.*
36: em_hvg_other.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	13	14.4	24	6	AX164353	AX164353 Sequence
2	13	14.4	26	6	AX039624	AX039624 Sequence
3	13	14.4	26	6	AX039654	AX039654 Sequence
4	13	14.4	30	6	AR028230	AR028230 Sequence
5	13	14.4	30	6	AR138633	AR138633 Sequence
6	13	14.4	33	5	XELARSE59	K01606 Xenopus lae
7	13	14.4	36	6	AX167671	AX167671 Sequence
8	13	14.4	39	6	AR011299	AR011299 Sequence
9	13	14.4	39	6	II17937	II17937 Sequence
10	12	13.3	14	6	AR082357	AR082357 Sequence
11	12	13.3	14	6	AR120899	AR120899 Sequence
12	12	13.3	14	6	I78403	I78403 Sequence
13	12	13.3	16	6	A14957	A14957 synthetic o
14	12	13.3	18	6	A14956	A14956 synthetic o
15	12	13.3	18	6	AR099365	AR099365 Sequence
16	12	13.3	20	6	A64909	A64909 Synthetic a
17	12	13.3	20	6	AR043539	AR043539 Sequence
18	12	13.3	21	6	AR063857	AR063857 Sequence
19	12	13.3	21	6	AR075821	AR075821 Sequence
20	12	13.3	21	6	AR098733	AR098733 Sequence
21	12	13.3	21	6	AR112325	AR112325 Sequence
22	12	13.3	21	6	E30454	E30454 Method for
23	12	13.3	21	6	E32364	E32364 Method for
24	12	13.3	21	6	E33635	E33635 Detection o
25	12	13.3	21	6	E33635	E33635 Detection o
26	12	13.3	21	6	E33635	E33635 Detection o
27	12	13.3	23	6	AR037890	AR037890 Sequence
28	12	13.3	24	6	AR054522	AR054522 Sequence
29	12	13.3	24	6	AR151501	AR151501 Sequence
30	12	13.3	25	6	AR082296	AR082296 Sequence
31	12	13.3	25	6	AR120838	AR120838 Sequence
32	12	13.3	25	6	AX115268	AX115268 Sequence
33	12	13.3	25	6	I78342	I78342 Sequence
34	12	13.3	27	6	AR060384	AR060384 Sequence
35	12	13.3	27	6	AR117878	AR117878 Sequence
36	12	13.3	27	6	AX114027	AX114027 Sequence
37	12	13.3	28	6	AR082955	AR082955 Sequence
38	12	13.3	28	6	AR082956	AR082956 Sequence
39	12	13.3	29	6	AX155821	AX155821 Sequence
40	12	13.3	30	6	A43687	A43687 Sequence
41	12	13.3	30	6	AR028176	AR028176 Sequence
42	12	13.3	30	6	AR050260	AR050260 Sequence
43	12	13.3	30	6	AR138579	AR138579 Sequence
44	12	13.3	30	6	AR140326	AR140326 Sequence
45	12	13.3	30	6	AX020976	AX020976 Sequence

ALIGNMENTS

RESULT 1
LOCUS AX164353 24 bp DNA
DEFINITION Sequence 183 from Patent WO0138564.
ACCESSION AX164353
VERSION AX164353.1 GI:14545287
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
REFERENCE 1 (bases 1 to 24)
AUTHORS Rouleau,G.A., Latremiere,R.G., Rochefort,D., Cossette,P. and
Ragsdale,D.
TITLE LocI for idiopathic generalized epilepsy, mutations thereof and
method using same to assess, diagnose, prognosis or treat epilepsy
JOURNAL Patent: WO 0138564-A 183 31-MAY-2001;
McGill University (CA)
FEATURES
source location/Qualifiers
1..24
/organism="synthetic construct"

/db_xref="taxon:32630"
BASE COUNT 2 a 5 c 3 g 14 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 34 ttatgtttcat 46
|||||
Db 8 TTTATGTTTCAT 20

RESULT 2
LOCUS AX039624 26 bp DNA PAT 18-NOV-2000
DEFINITION Sequence 13 from Patent WO0063441.
ACCESSION AX039624
VERSION AX039624.1 GI:11229653
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.
REFERENCE 1 (bases 1 to 26)
AUTHORS Herinstdt,C. and Davis,R.E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with alzheimer's disease
JOURNAL Patent: WO 0063441-A 13 26-OCT-2000;
MITOKOR (US)

FEATURES
Source 1. 26
/organism="synthetic construct"
/db_xref="taxon:32630"
BASE COUNT 8 a 2 c 5 g 11 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 37 atgtttcatgtt 49
|||||
Db 7 ATGTTTCATGTT 19

RESULT 3
LOCUS AX039654 26 bp DNA PAT 18-NOV-2000
DEFINITION Sequence 43 from Patent WO0063441.
ACCESSION AX039654
VERSION AX039654.1 GI:11229683
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.
REFERENCE 1 (bases 1 to 26)
AUTHORS Herinstdt,C. and Davis,R.E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with alzheimer's disease
JOURNAL Patent: WO 0063441-A 43 26-OCT-2000;
MITOKOR (US)

FEATURES
Source 1. 26
/organism="synthetic construct"
/db_xref="taxon:32630"
BASE COUNT 8 a 2 c 5 g 11 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 37 atgtttcatgtt 49
|||||
Db 7 ATGTTTCATGTT 19

RESULT 4
LOCUS AR028230 30 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 79 from patent US 5858661.
ACCESSION AR028230
VERSION AR028230.1 GI:5940203
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 30)
AUTHORS Shiloh,Y.
TITLE Ataxia-telangiectasia gene and its genomic organization
JOURNAL Patent: US 5858661-A 79 12-JAN-1999;
FEATURES
Source 1. 30
/organism="unknown"
BASE COUNT 13 a 2 c 1 g 14 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 aaaaattatttc 17
|||||
Db 15 AAAAATTATTTC 27

RESULT 5
LOCUS ARI38633 30 bp DNA PAT 16-JUN-2001
DEFINITION Sequence 158 from patent US 6200749.
ACCESSION ARI38633
VERSION ARI38633.1 GI:14480978
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 30)
AUTHORS Shiloh,Y.
TITLE Mutated forms of the ataxia-telangiectasia gene and method to screen for a partial A-T phenotype
JOURNAL Patent: US 6200749-A 158 13-MAR-2001;
FEATURES
Source 1. 30
/organism="unknown"
BASE COUNT 13 a 2 c 1 g 14 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 aaaaattatttc 17
|||||
Db 15 AAAAATTATTTC 27

RESULT 6
XELARSE59

LOCUS XELARSE59 33 bp DNA VRT 28-APR-1993
 DEFINITION Xenopus laevis autonomous replication sequence e59.
 ACCESSION K01606
 VERSION K01606.1 GI:213953
 KEYWORDS autonomous replication; mutational analysis.
 SOURCE Xenopus laevis DNA.
 ORGANISM Xenopus laevis
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Amphibia; Batrachia; Anura; Mesobatrachia; Pipidoidea; Pipidae;
 Xenopodinae; Xenopus.
 REFERENCE 1 (bases 1 to 33)
 AUTHORS Kearsey,S.
 TITLE Structural requirements for the function of a yeast chromosomal replicator
 JOURNAL Cell 37, 299-307 (1984)
 MEDLINE 84205653
 FEATURES
 source Location/Qualifiers
 1..33
 /organism="Xenopus laevis"
 /db_xref="taxon:8355"
 BASE COUNT 8 a 4 c 2 g 19 t
 ORIGIN

Query Match 14.4%; Score 13; DB 5; Length 33;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 31 atttattgtttt 43
 |||||||
 Db 1 ATTATTATGTTT 13

RESULT 7
 AX167671 36 bp DNA PAT 03-JUL-2001
 LOCUS AX167671
 DEFINITION Sequence 16 from Patent WO0144277.
 ACCESSION AX167671
 VERSION AX167671.1 GI:14597058
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM synthetic construct.
 REFERENCE 1 (bases 1 to 36)
 AUTHORS Wegrich Glover,L., Budziszewski,G.J., Levin,J.Z. and Zhou,Q.
 TITLE Herbicide target genes and methods
 JOURNAL Patent: WO 0144277-A 16 21-JUN-2001;
 Yngenta Participations AG (CH)
 FEATURES
 source Location/Qualifiers
 1..36
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="Oligonucleotide"
 BASE COUNT 6 a 4 c 11 g 15 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 36;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 36 tatgtttcatgt 48
 |||||||
 Db 21 TATGTTTCATCT 33

RESULT 8
 AR011299 39 bp DNA PAT 04-DEC-1998
 LOCUS AR011299/c
 DEFINITION Sequence 168 from patent US 5762938.
 ACCESSION AR011299
 VERSION AR011299.1 GI:3969289
 KEYWORDS

SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 39)
 AUTHORS Paolletti,E., Perkus,M.E., Taylor,J., Tartaglia,J., Norton,E.K.,
 Riviere,M., de Taisne,C., Limbach,K.J., Johnson,G.P., Pincus,S.E.,
 Cox,M.I., Audonnet,J.Francis and Gettig,R.Robert.
 TITLE Modified recombinant vaccinia virus and expression vectors thereof
 JOURNAL Patent: US 5762938-A 168 09-JUN-1998;
 FEATURES
 source Location/Qualifiers
 1..39
 /organism="unknown"
 BASE COUNT 15 a 6 c 7 g 11 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 39;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 ctgtaattttat 38
 |||||||
 Db 25 CTGTAATTTTAT 13

RESULT 9
 I17937 39 bp DNA PAT 07-OCT-1996
 LOCUS I17937/c
 DEFINITION Sequence 168 from patent US 5494807.
 ACCESSION I17937
 VERSION I17937.1 GI:1598292
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 39)
 AUTHORS Paolletti,E., Perkus,M.E., Taylor,J., Tartaglia,J., Norton,E.K.,
 Riviere,M., de Taisne,C., Limbach,K.J., Johnson,G.P., Pincus,S.E.,
 Cox,M.I., Audonnet,J.F. and Gettig,R.R.
 TITLE NYVAC Vaccinia virus recombinants comprising heterologous inserts
 JOURNAL Patent: US 5494807-A 168 27-FEB-1996;
 FEATURES
 source Location/Qualifiers
 1..39
 /organism="unknown"
 BASE COUNT 15 a 6 c 7 g 11 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 39;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 ctgtaattttat 38
 |||||||
 Db 25 CTGTAATTTTAT 13

RESULT 10
 AR082357 14 bp DNA PAT 31-AUG-2000
 LOCUS AR082357
 DEFINITION Sequence 201 from patent US 5972704.
 ACCESSION AR082357
 VERSION AR082357.1 GI:10009083
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 14)
 AUTHORS Draper,K.G., Chowrira,B., McSwigen,J., Stinchcomb,D.T. and
 Thompson,J.D.
 TITLE HIV nef targeted ribozymes
 JOURNAL Patent: US 5972704-A 201 26-OCT-1999;
 FEATURES Location/Qualifiers

source 1. .14
/organism="unknown"
BASE COUNT 7 a 3 c 3 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 11
LOCUS AR120899 14 bp DNA PAT 16-MAY-2001
DEFINITION Sequence 201 from patent US 6159692.
ACCESSION AR120899
VERSION AR120899.1 GI:14104475
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 14)
AUTHORS Draper,K.G., Chowrira,B., McSwiggen,J., Stinchcomb,D.T. and
TITLE Thompson,J.D.
METHOD Method and reagent for inhibiting human immunodeficiency virus
replication
JOURNAL Patent: US 6159692-A 201 12-DEC-2000;
FEATURES Location/Qualifiers
source 1. .14
BASE COUNT 7 a 3 c 3 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 12
LOCUS I78403 14 bp DNA PAT 03-APR-1998
DEFINITION Sequence 201 from patent US 5693535.
ACCESSION I78403
VERSION I78403.1 GI:3014557
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 14)
AUTHORS Draper,K.G., Chowrira,B., McSwiggen,J., Stinchcomb,D.T. and
TITLE Thompson,J.D.
METHOD HIV targeted ribozymes
JOURNAL Patent: US 5693535-A 201 02-DEC-1997;
FEATURES Location/Qualifiers
source 1. .14
BASE COUNT 7 a 3 c 3 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 13
LOCUS A14957 16 bp DNA PAT 18-FEB-1994
DEFINITION synthetic oligonucleotide (N2) from patent EP0211299.
ACCESSION A14957
VERSION A14957.1 GI:491869
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 16)
AUTHORS Habermann,P., Stengel,J.S. and Wengenmayer,F.
TITLE Fusion proteins, method for their production and their use
JOURNAL Patent: EP 0211299-A 2 25-FEB-1987;
FEATURES Location/Qualifiers
source 1. .16
BASE COUNT 1 a 4 c 2 g 9 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 16;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 15 CAATGAAGCAA 4

RESULT 14
LOCUS A14956 18 bp DNA PAT 18-FEB-1994
DEFINITION synthetic oligonucleotide (N1) from patent EP0211299.
ACCESSION A14956
VERSION A14956.1 GI:491868
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 18)
AUTHORS Habermann,P., Stengel,J.S. and Wengenmayer,F.
TITLE Fusion proteins, method for their production and their use
JOURNAL Patent: EP 0211299-A 1 25-FEB-1987;
FEATURES Location/Qualifiers
source 1. .18
BASE COUNT 9 a 3 c 5 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 4 CAATGAAGCAA 15

RESULT 15
LOCUS AR099365 18 bp DNA PAT 14-FEB-2001
DEFINITION Sequence 19 from patent US 6077709.

ACCESSION AR099365
VERSION AR099365.1 GI:12809131
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 18)
AUTHORS Bennett,C.Frank, Ackermann,E.J., Swayze,E.E. and Cowsett,L.M.
TITLE Antisense modulation of Survivin expression
JOURNAL Patent: US 6077709-A 19 20-JUN-2000;
FEATURES Location/Qualifiers
source 1..18
/organism="unknown"
BASE COUNT 1 a 3 c 3 g 11 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 51 tctattgttg 62
|||||
Db 1 TCTATTGTGG 12

RESULT 16
A07597 20 bp DNA PAT 09-JUL-1993
LOCUS A07597 Synthetic antisense oligonucleotide (5493-5512).
DEFINITION A07597
ACCESSION A07597.1 GI:413100
VERSION
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
REFERENCE 1 (bases 1 to 20)
ARTIFICIAL SEQUENCE.
AUTHORS Stropp,U., Baumgarten,J., Loebberding,A., Springer,W., Piel,N.,
Kretschmer,A., Koelbl,H. and Frommet,W.
TITLE Antisense-oligonucleotides for inhibiting the transactivator target
sequence (Tat) of HIV-1, and the synthesis of the transactivator protein
(Tat) of HIV-1, and their use
JOURNAL Patent: EP 0386563-A 4 12-SEP-1990;
ORIGIN BAYER AG

FEATURES Location/Qualifiers
source 1..20
/organism="synthetic construct"
BASE COUNT 8 a 4 c 3 g 5 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 74 caatgaagcaa 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 17
A64909 20 bp DNA PAT 29-MAR-1999
LOCUS A64909 Sequence 66 from Patent WO9731114.
DEFINITION A64909
ACCESSION A64909.1 GI:4530900
VERSION
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Burnham,M.K. and Hodgson,J.E.

TITLE POLYNUCLEOTIDES AND AMINOACID SEQUENCES FROM STAPHYLOCOCCUS AUREUS
JOURNAL Patent: WO 9731114-A 66 28-AUG-1997;
SMITHKLINE BEECHAM PLC (GB)
FEATURES Location/Qualifiers
source 1..20
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 11 a 4 c 2 g 3 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
Db 20 GTTTCACGTTT 9

RESULT 18
AR043539 21 bp DNA PAT 29-SEP-1999
LOCUS AR043539
DEFINITION Sequence 6 from patent US 5814490.
ACCESSION AR043539
VERSION AR043539.1 GI:5964547
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Spears,P.A.
TITLE Amplification and detection of chlamydia trachomatis nucleic acids
JOURNAL Patent: US 5814490-A 6 29-SEP-1998;
FEATURES Location/Qualifiers
source 1..21
/organism="unknown"
BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
Db 13 GTTTCACGTTT 2

RESULT 19
AR063857 21 bp DNA PAT 29-SEP-1999
LOCUS AR063857
DEFINITION Sequence 4 from patent US 5846726.
ACCESSION AR063857
VERSION AR063857.1 GI:5993165
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Nadeau,J.G., Pither,J.Bruce, Schram,J.L., Linn,C.Preston, Vonk,G.P.
TITLE Detection of nucleic acids by fluorescence quenching
JOURNAL Patent: US 5846726-A 4 08-DEC-1998;
FEATURES Location/Qualifiers
source 1..21
/organism="unknown"
BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;

Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGTTT 2

RESULT 20
LOCUS AR075821/c 21 bp DNA PAT 30-AUG-2000

DEFINITION Sequence 4 from patent US 5958700.

ACCESSION AR075821

VERSION AR075821.1 GI:10002567

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 21)

AUTHORS Nadeau,J.G., Pitner,J.Bruce, Linn,C.Preston and Schram,J.L.

TITLE Detection of nucleic acids by fluorescence quenching

JOURNAL Patent: US 5958700-A 4 28-SEP-1999;

FEATURES Location/Qualifiers

source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGTTT 2

RESULT 21

LOCUS AR098733/c 21 bp DNA PAT 14-FEB-2001

DEFINITION Sequence 8 from patent US 6077669.

ACCESSION AR098733

VERSION AR098733.1 GI:12808499

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 21)

AUTHORS Little,M.C. and Vonk,G.P.

TITLE Kit and method for fluorescence based detection assay

JOURNAL Patent: US 6077669-A 8 20-JUN-2000;

FEATURES Location/Qualifiers

source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGTTT 2

RESULT 22

LOCUS AR112325/c 21 bp DNA PAT 16-MAY-2001

DEFINITION Sequence 5 from patent US 6130047.

ACCESSION AR112325

VERSION AR112325.1 GI:14092225

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 21)

AUTHORS Nadeau,J.G., Hsieh,H.V., Pitner,J.Bruce and Linn,C.Preston.

TITLE Detection of nucleic acids by fluorescence quenching

JOURNAL Patent: US 6130047-A 5 10-OCT-2000;

FEATURES Location/Qualifiers

source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGTTT 2

RESULT 23

LOCUS E30454/c 21 bp DNA PAT 07-FEB-2001

DEFINITION Method for detecting target nucleic acid sequence and

oligonucleotide.

ACCESSION E30454

VERSION E30454.1 GI:13025611

KEYWORDS JP 1999056380-A/4.

SOURCE unidentified.

ORGANISM unidentified.

REFERENCE 1 (bases 1 to 21)

AUTHORS James,G.N.J., Pitona,C.P.R.R. and L,S.

TITLE Method for detecting target nucleic acid sequence and

JOURNAL Patent: JP 1999056380-A 4 02-MAR-1999;

COMMENT BECTON DICKINSON & CO

OS Unidentified

PN JP 1999056380-A/4

PD 02-MAR-1999

PE 29-MAY-1998 JP 1998166141

PR 30-MAY-1997 US 08/865 675

PI JAMES G NADEAU, J BRUCE PITONA, C PRESTON RIN, JAMES L SHURAMU PC

CI2N15/09,CI2Q1/68,G01N33/50,G01N33/366,CI2N15/00 CC

Strandedness: Single;

CC Topology: Linear;

EH Key

FT source 1..21

FEATURES Location/Qualifiers

source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGTTT 2

RESULT 24

LOCUS E32364/c 21 bp DNA PAT 07-FEB-2001

ACCESSION E32364

```

DEFINITION Method for detecting nucleic acid by fluorescent quenching.
ACCESSION E33364
VERSION E33364.1 GI:13026696
KEYWORDS JP 1999123083-A/4.
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 21)
AUTHORS James,G.N.J., Pitona,J.L.S., Preston,R.G.P. and G.T.W.
TITLE Method for detecting nucleic acid by fluorescent quenching
JOURNAL Patent: JP 1999123083-A 4 11-MAY-1999;
BECTON DICKINSON & CO
COMMENT OS Unidentified
PN JP 1999123083-A/4
PD 11-MAY-1999
PF 13-MAY-1997 US 08/855 085
PI JAMES G. MADEAU, J. BLUCE PITONA, JAMES L. SHURAMU, C. PRESTON RIN,
PI GREN P VONG,
PI G. TERANSU WALKER
PC C12N15/09, C07H21/00, C12Q1/68, G01N21/64, G01N33/58, C12N15/00 CC
Strandedness: Single;
CC Topology: Linear;
FH Key Location/Qualifiers
FT source 1..21
/organism='Unidentified'.
FEATURES
SOURCE Location/Qualifiers
1..21
/organism='unidentified'
/db_xref='taxon:32644'
BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 39 gtttcacgttt 50
| | | | | | | | | | | | | | | | | | | | |
Db 13 GTTTCACGTTT 2

RESULT 25
E33635/c 21 bp DNA PAT 07-FEB-2001
DEFINITION Detection of nucleic acid by disappearance of fluorescence.
ACCESSION E33635
VERSION E33635.1 GI:13027030
KEYWORDS JP 1999155598-A/5.
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 21)
AUTHORS James,G.N.H., C.J.B.P.P. and Preston, R.
TITLE Detection of nucleic acid by disappearance of fluorescence
JOURNAL Patent: JP 1999155598-A 5 15-JUN-1999;
BECTON DICKINSON & CO
COMMENT OS Artificial Sequence
PN JP 1999155598-A/5
PD 15-JUN-1999
PF 22-SEP-1998 JP 1998267492
PR 23-SEP-1997 US 08/933749
PI JAMES G. MADEAU, HELEN V C, J. BLUCE PITONA, C. PRESTON RIN PC
C12Q1/68, C07H21/00, C12N15/09, G01N21/64, G01N33/542, G01N33/566, PC
C12N15/00
CC
FH Key Location/Qualifiers
FT source 1..21
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FEATURES
SOURCE Location/Qualifiers
1..21
/organism='unidentified'

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BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 39 gtttcacgttt 50
| | | | | | | | | | | | | | | | | | | | |
Db 13 GTTTCACGTTT 2

RESULT 27
AR037890/c 23 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 10 from patent US 5804383.
ACCESSION AR037890
VERSION AR037890.1 GI:5956607
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 23)
AUTHORS Grenert,D.C. and Dohman,A.F.
TITLE Method and assay for detection of the expression of allele-specific
polymers by allele-specific in situ reverse transcriptase
JOURNAL Patent: US 5804383-A 10 08-SEP-1998;
FEATURES
SOURCE Location/Qualifiers
1..21
/organism='unidentified'
/db_xref='taxon:32644'
BASE COUNT 8 a 5 c 3 g 5 t.
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 39 gtttcacgttt 50
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Db 13 GTTTCACGTTT 2

FEATURES
SOURCE Location/Qualifiers
1..21
/organism='Artificial Sequence'.

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source 1. .23
/organism="unknown"
BASE COUNT 4 a 4 c 3 g 12 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 23;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 59 ttggagcaataa 70
|||||
Db 23 TTGGAGCAATAA 12

RESULT 28
AR054522 AR054522 24 bp DNA PAT 29-SEP-1999
LOCUS Sequence 31 from patent US 5837441.
DEFINITION AR054522
ACCESSION AR054522
VERSION AR054522.1 GI:5980099
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 24)
AUTHORS Hjelte, B. and Jensen, S.
TITLE Hantavirus-associated respiratory distress virus antigens
JOURNAL Patent: US 5837441-A 31 17-NOV-1998;
FEATURES Location/Qualifiers
source 1. .24
/organism="unknown"

BASE COUNT 8 a 1 c 7 g 8 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 21 gtttactgtaac 32
|||||
Db 2 GTTACTGTAAT 13

RESULT 29
AR151501 AR151501 24 bp DNA PAT 08-AUG-2001
LOCUS Sequence 26 from patent US 6232094.
DEFINITION AR151501
ACCESSION AR151501
VERSION AR151501.1 GI:15117551
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 24)
AUTHORS Hanson, L., Stromqvist, M., Bergstrom, S., Hernell, O., and Tornell, J.
TITLE DNA encoding human kappa, casein and process for obtaining the protein
JOURNAL Patent: US 6232094-A 26 15-MAY-2001;
FEATURES Location/Qualifiers
source 1. .24
/organism="unknown"
BASE COUNT 5 a 3 c 2 g 14 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 35 ttaatgttcaat 46
|||||

Db 12 TTATGTTTCAAT 23

RESULT 30
AR082296/C AR082296 25 bp DNA PAT 31-AUG-2000
LOCUS Sequence 140 from patent US 5972704.
DEFINITION AR082296
ACCESSION AR082296
VERSION AR082296.1 GI:10009022
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Draper, K.G., Chowrira, B., McSwiggen, J., Stinchcomb, D.T. and
TITLE HIV net targeted ribozymes
JOURNAL Patent: US 5972704-A 140 26-OCT-1999;
FEATURES Location/Qualifiers
source 1. .25
/organism="unknown"

BASE COUNT 6 a 4 c 5 g 10 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 74 caatgaagcaaa 85
|||||
Db 23 CAATGAAGCAA 12

RESULT 31
AR120838/C AR120838 25 bp DNA PAT 16-MAY-2001
LOCUS Sequence 140 from patent US 6159692.
DEFINITION AR120838
ACCESSION AR120838
VERSION AR120838.1 GI:14104414
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Draper, K.G., Chowrira, B., McSwiggen, J., Stinchcomb, D.T. and
TITLE Method and reagent for inhibiting human immunodeficiency virus replication
JOURNAL Patent: US 6159692-A 140 12-DEC-2000;
FEATURES Location/Qualifiers
source 1. .25
/organism="unknown"

BASE COUNT 6 a 4 c 5 g 10 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 74 caatgaagcaaa 85
|||||
Db 23 CAATGAAGCAA 12

RESULT 32
AX115268 AX115268 25 bp DNA PAT 11-MAY-2001
LOCUS Sequence 391 from Patent WO0129262.
DEFINITION AX115268
ACCESSION AX115268
VERSION AX115268.1 GI:14032210
KEYWORDS

SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequence.
REFERENCE 1 (bases 1 to 25)
AUTHORS Picoult-Newburg, L. and Pohl, M.
TITLE Genotyping reagents, kits and methods of use thereof
JOURNAL Patent: WO 0129262-A 391 26-APR-2001;
Orchid Biosciences, Inc. (US)
FEATURES Location/Qualifiers
SOURCE 1..25
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Primer"
BASE COUNT 8 a 4 c 3 g 10 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 7 aaattattca 18
DB 3 AAAATTATTCA 14
RESULT 33
178342/c 178342 25 bp DNA PAT 03-APR-1998
LOCUS Sequence 140 from patent US 5693535.
DEFINITION 178342
ACCESSION 178342
VERSION 178342.1 GI:3014496
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Draper, K.G., Chowitra, B., McSwiggen, J., Stinchcomb, D.T. and
Thompson, J.D.
TITLE HIV targeted ribozymes
JOURNAL Patent: US 5693535-A 140 02-DEC-1997;
FEATURES Location/Qualifiers
SOURCE 1..25
/organism="unknown"
BASE COUNT 6 a 4 c 5 g 10 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 74 caatgaagcaaa 85
DB 23 CAATGAAGCAA 12
RESULT 34
AR060384/c 27 bp DNA PAT 29-SEP-1999
LOCUS AR060384
DEFINITION Sequence 6 from patent US 5840568.
ACCESSION AR060384
VERSION AR060384.1 GI:5986834
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Pfeundschuh, M.
TITLE Hodgkin's disease associated molecules and uses thereof
JOURNAL Patent: US 5840568-A 6 24-NOV-1998;
FEATURES Location/Qualifiers
SOURCE 1..27

BASE COUNT 5 a 7 c 4 g 11 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 77 tgaagcaagt 88
DB 26 TGAAGCAAGTG 15
RESULT 35
AR117878/c 27 bp DNA PAT 16-MAY-2001
LOCUS AR117878
DEFINITION Sequence 6 from patent US 6140464.
ACCESSION AR117878
VERSION AR117878.1 GI:14098784
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Pfeundschuh, M. and Rammensee, H.
TITLE Nonapeptides that bind a HLA-A2.1 molecule
JOURNAL Patent: US 6140464-A 6 31-OCT-2000;
FEATURES Location/Qualifiers
SOURCE 1..27
/organism="unknown"
BASE COUNT 5 a 7 c 4 g 11 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 77 tgaagcaagt 88
DB 26 TGAAGCAAGTG 15
RESULT 36
AX114027/c 27 bp DNA PAT 08-MAY-2001
LOCUS AX114027
DEFINITION Sequence 6 from Patent EP1108432.
ACCESSION AX114027
VERSION AX114027.1 GI:14018204
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 27)
AUTHORS Pfeundschuh, M.I.
TITLE Method for identifying or isolating a molecule and molecules
JOURNAL identified thereby
JOURNAL Patent: EP 1108432-A 6 20-JUN-2001;
FEATURES LUDWIG INSTITUTE FOR CANCER RESEARCH (US)
SOURCE 1..27
Location/Qualifiers
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 5 a 7 c 4 g 11 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 77 tgaagcaagt 88

Db 26 TGAAGCAAGTG 15

RESULT 37

AR082955 AR082955 28 bp DNA PAT 01-SEP-2000
 LOCUS Sequence 20 from patent US 5976795.
 DEFINITION
 AR082955
 ACCESSION AR082955.1 GI:10009745
 VERSION
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 28)
 AUTHORS Voytas,D.F. and Zou,S.
 TITLE Retrotransposon and methods
 JOURNAL Patent: US 5976795-A 20 02-NOV-1999;
 FEATURES Location/Qualifiers
 source 1..28

BASE COUNT 7 a 0 c 4 g 15 t 2 others
 ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 28;
 Best Local Similarity 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 31 attttatgtt 42
 Db 7 ATTTTATGTTT 18

RESULT 38
 AR082956/C AR082956 28 bp DNA PAT 01-SEP-2000
 LOCUS Sequence 21 from patent US 5976795.
 DEFINITION
 AR082956
 ACCESSION AR082956
 VERSION AR082956.1 GI:10009746
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 28)
 AUTHORS Voytas,D.F. and Zou,S.
 TITLE Retrotransposon and methods
 JOURNAL Patent: US 5976795-A 21 02-NOV-1999;
 FEATURES Location/Qualifiers
 source 1..28

BASE COUNT 15 a 4 c 0 g 7 t 2 others
 ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 28;
 Best Local Similarity 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 31 attttatgtt 42
 Db 22 ATTTTATGTTT 11

RESULT 39
 AX155921/C AX155921 29 bp DNA PAT 22-JUN-2001
 LOCUS Sequence 164 from patent WO0140474.
 DEFINITION
 AX155921
 ACCESSION AX155921.1 GI:14537028
 VERSION
 KEYWORDS
 SOURCE Chlamydia sp.
 ORGANISM Chlamydia sp.

REFERENCE 1 (bases 1 to 29)
 AUTHORS Probst,P., Bhatia,A., Skeiky,Y.A., Flinn,S.P. and Scholler,J.
 TITLE Compounds and methods for treatment and diagnosis of chlamydial
 JOURNAL Infection
 Patent: WO 0140474-A 164 07-JUN-2001;
 FEATURES CORIXA CORPORATION (US)
 source 1..29

BASE COUNT 12 a 4 c 4 g 9 t
 ORIGIN /db_xref="taxon:35827"

Query Match 13.3%; Score 12; DB 6; Length 29;
 Best Local Similarity 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 48 ttctctatgt 59
 Db 15 TTTCTATTGT 4

RESULT 40
 A43687/C A43687 30 bp DNA PAT 06-MAR-1997
 LOCUS Sequence 3 from Patent WO9508642.
 DEFINITION
 A43687
 ACCESSION A43687
 VERSION A43687.1 GI:2298875
 KEYWORDS
 SOURCE unidentified.
 ORGANISM unidentified.

REFERENCE 1 (bases 1 to 30)
 AUTHORS Sammes,P.G. and Garman,A.J.
 TITLE NUCLEIC ACID DETECTION WITH ENERGY TRANSFER
 JOURNAL Patent: WO 9508642-A 3 30-MAR-1995;
 ZENECA LTD (GB)

COMMENT Other publication AU 7662794 950410
 Other publication GB 2283095 950426.
 FEATURES Location/Qualifiers
 source 1..30

BASE COUNT 12 a 4 c 4 g 10 t
 ORIGIN /db_xref="taxon:32644"

Query Match 13.3%; Score 12; DB 6; Length 30;
 Best Local Similarity 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 26 ctgtaatttta 37
 Db 29 CTGTATTTTTA 18

RESULT 41
 AR028176/C AR028176 30 bp DNA PAT 29-SEP-1999
 LOCUS Sequence 25 from patent US 5858661.
 DEFINITION
 AR028176
 ACCESSION AR028176
 VERSION AR028176.1 GI:5940149
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
 AUTHORS Shilon,Y.
 TITLE Ataxia-telangiectasia gene and its genomic organization
 JOURNAL Patent: US 5858661-A 25 12-JAN-1999;
 FEATURES Location/Qualifiers

source 1..30
/organism="unknown"
BASE COUNT 8 a 2 c 2 g 18 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaattatt 15
|||||
Db 19 AAAAAATTATT 8

RESULT 42
AR050260/c 30 bp DNA PAT 29-SEP-1999
LOCUS
DEFINITION Sequence 3 from patent US 5827653.
ACCESSION AR050260
VERSION AR050260.1 GI:5972985
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
AUTHORS Sammes,P.George and Garman,A.John.
TITLE Nucleic acid detection with energy transfer
JOURNAL Patent: US 5827653-A 3 27-OCT-1998;
FEATURES Location/Qualifiers
source 1..30
/organism="unknown"

BASE COUNT 12 a 4 c 4 g 10 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 ctgtaattttta 37
|||||
Db 29 CTGTAATTTTAA 18

RESULT 43
ARI38579/c 30 bp DNA PAT 16-JUN-2001
LOCUS
DEFINITION Sequence 104 from patent US 6200749.
ACCESSION ARI38579
VERSION ARI38579.1 GI:14480924
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
AUTHORS Shiloh,Y.
TITLE Mutated forms of the ataxia-telangiectasia gene and method to
screen for a partial A-T phenotype
JOURNAL Patent: US 6200749-A 104 13-MAR-2001;
FEATURES Location/Qualifiers
source 1..30
/organism="unknown"

BASE COUNT 8 a 2 c 2 g 18 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaattatt 15
|||||

Db 19 AAAAAATTATT 8

RESULT 44
ARI40326/c 30 bp DNA PAT 16-JUN-2001
LOCUS
DEFINITION Sequence 14 from patent US 6207455.
ACCESSION ARI40326
VERSION ARI40326.1 GI:14482822
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
AUTHORS Chang,L.-J.
TITLE Lentiviral vectors
JOURNAL Patent: US 6207455-A 14 27-MAR-2001;
FEATURES Location/Qualifiers
source 1..30
/organism="unknown"

BASE COUNT 7 a 5 c 6 g 12 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagcaaa 85
|||||
Db 28 CAATGAAGCAAA 17

RESULT 45
AX020976/c 30 bp DNA PAT 07-SEP-2000
LOCUS
DEFINITION Sequence 22 from Patent EP0928832.
ACCESSION AX020976
VERSION AX020976.1 GI:10044639
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequence.

REFERENCE 1 (bases 1 to 30)
AUTHORS Kelly,S.J., Weston,S.L. and Robertson,N.H.
TITLE Cyclic fibrosis test based on the detection of mutations in the
cfr gene by arms
JOURNAL Patent: EP 0928832-A 22 14-JUL-1999;
FEATURES ZENECA LTD (GB)
source Location/Qualifiers
1..30
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR Amplification Primer"

BASE COUNT 8 a 6 c 5 g 11 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 59 ttgaggaataa 70
|||||
Db 17 TTGAGGACATAA 6

Search completed: January 24, 2002, 03:22:38
Job time: 3815 sec

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BASE COUNT 2 a 5 c 3 g 14 t
 ORIGIN /db_xref="taxon:32630"
 /note="synthetic oligonucleotide"

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 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 34 ttatgtttcat 46
 Db 8 TTTATGTTTCAT 20

RESULT 2
 AX039624 26 bp DNA PAT 18-NOV-2000
 LOCUS AX039624
 DEFINITION Sequence 13 from Patent WO0063441.
 ACCESSION AX039624
 VERSION AX039624.1 GI:11229653
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM artificial sequence.
 REFERENCE 1 (bases 1 to 26)
 AUTHORS Herinstad,C. and Davis,R.E.
 TITLE Single nucleotide polymorphisms in mitochondrial genes that segreg
 JOURNAL Patent: WO 0063441-A 13 26-OCT-2000;
 MITOKOR (US)

FEATURES
 source Location/Qualifiers
 1..26
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="PCR primer"
 BASE COUNT 8 a 2 c 5 g 11 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 26;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 37 atgttcattgt 49
 Db 7 ATGTTTCATGTT 19

RESULT 3
 AX039654 26 bp DNA PAT 18-NOV-2000
 LOCUS AX039654
 DEFINITION Sequence 43 from Patent WO0063441.
 ACCESSION AX039654
 VERSION AX039654.1 GI:11229663
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM artificial sequence.
 REFERENCE 1 (bases 1 to 26)
 AUTHORS Herinstad,C. and Davis,R.E.
 TITLE Single nucleotide polymorphisms in mitochondrial genes that segreg
 JOURNAL Patent: WO 0063441-A 43 26-OCT-2000;
 MITOKOR (US)

FEATURES
 source Location/Qualifiers
 1..26
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="PCR primer"

BASE COUNT 8 a 2 c 5 g 11 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 26;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 atgttcattgt 49
 Db 7 ATGTTTCATGTT 19

RESULT 4
 AR028230 30 bp DNA PAT 29-SEP-1999
 LOCUS AR028230
 DEFINITION Sequence 79 from patent US 5858661.
 ACCESSION AR028230
 VERSION AR028230.1 GI:5940203
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 30)
 AUTHORS Shiloh,Y.
 TITLE Ataxia-telangiectasia gene and its genomic organization
 JOURNAL Patent: US 5858661-A 79 12-JAN-1999;
 FEATURES Location/Qualifiers
 source 1..30
 /organism="unknown"

BASE COUNT 13 a 2 c 1 g 14 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 5 aaaaattatttc 17
 Db 15 AAAAATTATTTC 27

RESULT 5
 ARI38633 30 bp DNA PAT 16-JUN-2001
 LOCUS ARI38633
 DEFINITION Sequence 158 from patent US 6200749.
 ACCESSION ARI38633
 VERSION ARI38633.1 GI:14480978
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 30)
 AUTHORS Shiloh,Y.
 TITLE Mutated forms of the ataxia-telangiectasia gene and method to
 screen for a partial A-T phenotype
 JOURNAL Patent: US 6200749-A 158 13-MAR-2001;
 FEATURES Location/Qualifiers
 source 1..30
 /organism="unknown"

BASE COUNT 13 a 2 c 1 g 14 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 aaaaattatttc 17
 Db 15 AAAAATTATTTC 27

RESULT 6
 XELARSE59

Query Match 4.9%; Score 16; DB 20; Length 32;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 189 aaataattattttt 204
Db 4 aaataattattttt 19
|||||

RESULT 4
AAZ97603
ID AAT97603 standard; DNA: 36 BP.
XX
AC AAT97603;
AT 30-APR-1998 (first entry)

XX Shigella dysenteriae delta-stx-A allele PCR primer 13.
XX Delta-virG allele; delta-guaB-A allele; PCR: amplification; primer;
KW delta-stx-A allele; shigellosis; vaccine; ss.
XX Synthetic.
OS Shigella dysenteriae.

XX WO9737685-A1.

XX PN 16-OCT-1997.

XX PD 09-APR-1997; 97WO-US05954.

XX PF 09-APR-1996; 96US-0629600.

XX PR (UWMA-) UNIV MARYLAND BALTIMORE.

XX Levine MM, Noriega FR;
PI WPI: 1997-512417/47.

XX Shigella mutants with mutation in guaB-A - used in vaccines against
XX Shigellosis

XX Example 6; Page 57; 94pp; English.

XX This is a PCR primer used in the amplification of the Shigella
XX dysenteriae 1 delta-stx-A allele. The delta-stx-A allele was integrated
XX into delta-guaB-A of delta-guaB-A, delta-virG S. dysenteriae 1, which
XX inactivated the shiga toxin of this strain. The mutant can be used in
XX the preparation of vaccines such as, a live vector vaccine comprising
XX a Shigella mutant, (which encodes and expresses a foreign antigen, and
XX a pharmaceutically acceptable carrier) or a DNA mediated vaccine
XX comprising the Shigella mutant (which also contains a plasmid which
XX encodes and expresses a foreign antigen in a eukaryotic cell). The
XX vaccines can be used against Shigellosis.

XX Sequence 36 BP; 11 A; 3 C; 10 G; 12 T; 0 other;

Query Match 4.9%; Score 16; DB 18; Length 36;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Yy 189 aaataattattttt 204
Db 14 aaataattattttt 29
|||||

RESULT 5
AAV90775
D AAV90775 standard; DNA: 39 BP.

AC AAV90775;
XX 18-FEB-1999 (first entry)
XX Primer Y104F.
XX Antigen; immunogenic cluster family; vaccine; gastritis; diagnosis;
KW peptic ulcer; gastric adenocarcinoma; gastric lymphoma; primer; ss.
XX Synthetic.
XX WO9849314-A2.
XX PD 05-NOV-1998.
XX PF 27-APR-1998; 98WO-US08487.
XX PR 14-OCT-1997; 97US-0061958.
XX PR 25-APR-1997; 97US-0045107.
XX (GENE-) GENELABS TECHNOLOGIES INC.
XX Chow TP, Fry KE, Lim MY, McAtee CP;
XX WPI: 1999-009433/01.
XX New Helicobacter pylori antigens and related nucleic acid sequences
XX - useful in serological diagnosis and protective vaccines, providing
XX long-lasting immune response
XX Claim Disclosure; Page 194; 402pp; English.
XX The specification, which describes Helicobacter pylori antigenic
XX proteins that are characterised by immunoreactivity with
XX H. pylori-positive antisera. The specification also describes 69
XX previously unrecognised immunogenic cluster families. H. pylori
XX antigens are used to detect H. pylori-specific antibodies, for
XX diagnosing infection or to confirm eradication of infection, and
XX in vaccines to protect against H. pylori infection and related
XX diseases (gastritis, peptic ulcer, gastric adenocarcinoma/lymphoma).
XX The present primer is used in the course of the invention.
XX Sequence 39 BP; 15 A; 8 C; 7 G; 9 T; 0 other;

Query Match 4.9%; Score 16; DB 20; Length 39;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 atattcttaatttagc 23
Db 23 atattcttaatttagc 38
|||||

RESULT 6
AAZ68379
ID AAZ68379 standard; DNA: 47 BP.
XX AAZ68379;

XX 10-SEP-2001 (first entry)
XX Human map-related biallelic marker SEQ ID NO:2726.

XX Human genome; biallelic marker; high density disequilibrium map;
XX genomic map; haplotype; phenotype; polymorphic base; genotyping;
XX haplotyping; hybridisation; identification; characterisation;
XX diagnosis; single nucleotide polymorphism; SNP; ds.
XX Homo sapiens.

XX Key Location/Qualifiers
XX Variation replace(24,A)

SECRET
AC AAV9
XX
DT 18-F
XX

30-APR-1998 (first entry)

Shigella dysenteriae delta-stxA allele PCR primer 13

Delta-virc allele; delta-guab-A allele; PCR; amplification; primer

delta-stix alliere; singeliosos; vacuone; z

Synthetic.

XX

MOJ/3/000 HT
PN
XX

PD 16-OCT-1997

PF 09-APR-1997: 97WO-US05954

09-APR-1996; 96US-0629600

UNIV MARYLAND BALTIMORE.

Levine MM, Noriega FR;

WPI; 1997-512417/47.

Shigella mutants with mutation in *guab-A* - used in vaccines against Shigellosis

Example 6; Page 57; 94pp; English

xx This is a PCR primer used in the amplification of the Shigella
 cc dysenteriae 1 delta-stx allele. The delta-stx allele was integrated
 cf into delta-guab-A of delta-guab-A, delta-virid S. dysenteriae 1, which
 cd inactivated the shiga toxin of this strain. The mutant can be used in
 cc the preparation of vaccines such as, a live vector vaccine comprising
 cf a Shigella mutant, (which encodes and expresses a foreign antigen, and
 cc a pharmaceutically acceptable carrier) or a DNA mediated vaccine
 cc comprising the Shigella mutant (which also contains a plasmid which
 cc encodes and expresses a foreign antigen in a eukaryotic cell). The
 cc vaccines can be used against Shigellosis.
 'xx
 50 Sequence 36 BP; 11 A; 3 C; 10 G; 12 T; 0 other;

Sequence 36 BP; 11 A; 3 C; 10 G; 12 T; 0 other

```

Query Match          4.98; Score 16; DB 18; Length 36;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0.

```

```

QY      189 aaataattatttta 204
        |||||
Db      14 aaataattatttta 29

```

RESULT	5
AAV90775	
ID	AAV90775 standard; DNN: 39 BP.
XX	

AC AAV91

AC AAV90775;

DT 18-FEB-1999 (first entry)

DE Primer Y104F

XX Antigen; ImmH₁ cluster family; vaccine; gastritis; diagnosis
KW Gastric adenocarcinoma; gastric lymphoma; primer; ss
KM Peptic ulcer; gastric adenocarcinoma; gastric lymphoma; primer; ss

Synthetic

PN WO9849314-A2
XX

02-NOV-1990
PD
XX

XX

PR 25-APR-1997; 97US-0045107

PA (GENE-) GENELABS TECHNOLOGIES INC.

PI Chow TP, Fry KE, Lill MI, Maceee C

DR WPL: 1999-003433/01
XY

PT - useful in serological diagnosis and protective vaccines, providing

XX

ps Claim Disclosure; page 194; 402pp; English

XX The specification, which describes *Helicobacter pylori* antigenic
CC proteins that are characterised by immunoreactivity with
CC H. pylori-positive antisera. The specification also describes 69
CC H. pylori-positive antisera. The specification also describes 69
CC previously unrecognised immunogenic cluster families. H. pylori
CC antigens are used to detect H. pylori-specific antibodies, for
CC diagnosing infection or to confirm eradication of infection, and
CC in vaccines to protect against H. pylori infection and related
CC diseases (gastritis, peptic ulcer, gastric adenocarcinoma/Lymphoma)
CC The present primer is used in the course of the invention.
XX
CC
SO Sequence 39 BP: 15 A; 8 C; 7 G; 9 T; 0 other:

Sequence 39 BP; 12 A, 0 C, 7 G, 2 T, 2 TTTT

Query Match	4.9%	Score 16	DB 20	Length 39
-Best Local Similarity	.100.0%	Pred. No. 1e+03		
Matches 16	Conservative 0	Mismatches 0	Indels 0	Gaps 0

8 atattctaaat.tltagc 23
QY

Db 23 atatcttaatttagc 3b

RESULT	6
AA268379	
ID	AA268379 standard; DNA: 47 BP.

AC AAZ68379;

DT 10-SEP-2001 (FIRST ENTRY)
 YY

DE Human map-related diallelic marker SED ID NO: 2/20

XX	Human genome; biallelic marker; high density disequilibrium map;
XX	
KM	genomic map; haplotype; phenotype; polymorphic base; genotyping;
KM	haplotyping; hybridisation; identification; characterisation;
KM	diagnosis; single nucleotide polymorphism; SNP; ds.
XY	

OS Homo sapiens

	Key	Location/Qualifiers
FH	variation	replace(24,A)
FT		

us-09-531-438-3.oli.rng

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:27:08 ; Search time 222.28 seconds

Title: US-09-531-438-3

Perfect score: 327
Sequence: 1 atttgggatatctaaattt.....tttcatgttttcttattgtt 327

Scoring table: OLIGO_NUC

Searched: 930621.seqs, 428662619 residues

Word size : 0

Total number of hits satisfying chosen parameters: 989696

Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database : N_Geneseq_1101: *

1.	/SID52/cgcgdata/geneseq/geneseqn/NA1980.DAT.*
2.	/SID52/cgcgdata/geneseq/geneseqn/NA1981.DAT.*
3.	/SID52/cgcgdata/geneseq/geneseqn/NA1982.DAT.*
4.	/SID52/cgcgdata/geneseq/geneseqn/NA1983.DAT.*
5.	/SID52/cgcgdata/geneseq/geneseqn/NA1984.DAT.*
6.	/SID52/cgcgdata/geneseq/geneseqn/NA1985.DAT.*
7.	/SID52/cgcgdata/geneseq/geneseqn/NA1986.DAT.*
8.	/SID52/cgcgdata/geneseq/geneseqn/NA1987.DAT.*
9.	/SID52/cgcgdata/geneseq/geneseqn/NA1988.DAT.*
10.	/SID52/cgcgdata/geneseq/geneseqn/NA1989.DAT.*
11.	/SID52/cgcgdata/geneseq/geneseqn/NA1990.DAT.*
12.	/SID52/cgcgdata/geneseq/geneseqn/NA1991.DAT.*
13.	/SID52/cgcgdata/geneseq/geneseqn/NA1992.DAT.*
14.	/SID52/cgcgdata/geneseq/geneseqn/NA1993.DAT.*
15.	/SID52/cgcgdata/geneseq/geneseqn/NA1994.DAT.*
16.	/SID52/cgcgdata/geneseq/geneseqn/NA1995.DAT.*
17.	/SID52/cgcgdata/geneseq/geneseqn/NA1996.DAT.*
18.	/SID52/cgcgdata/geneseq/geneseqn/NA1997.DAT.*
19.	/SID52/cgcgdata/geneseq/geneseqn/NA1998.DAT.*
20.	/SID52/cgcgdata/geneseq/geneseqn/NA1999.DAT.*
21.	/SID52/cgcgdata/geneseq/geneseqn/NA2000.DAT.*
22.	/SID52/cgcgdata/geneseq/geneseqn/NA2001.DAT.*

Pred. No.' is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Result No.	Score	Query Match	Length	DB ID	Description
1	16	4.9	29	AA742655	Primer for amplify
2	16	4.9	29	AA551200	N-terminal primer
3	16	4.9	32	AA327689	PCR primer for Ver
4	16	4.9	36	AA797603	Shigella dysenteriae
5	16	4.9	39	AA797603	Primer Y104F. Syn
6	16	4.9	47	AA268379	Human map-related
7	15	4.6	20	AA466183	Dog genomic marker
8	15	4.6	24	AA446804	Human high motilit
9	15	4.6	27	AA739439	Hel-N2 selected se
10	15	4.6	27	AAV37457	Human Hel-N2 selec
11	15	4.6	30	AA043975	Triple helix formi

12	15	4	6	48	16	AAT35576	Human gene signatu
13	14	4	3	18	13	AAQ20161	Cross-linking olig
14	14	4	3	18	13	AAQ20160	Cross-linking olig
15	14	4	3	18	13	AAQ30311	Oligomer HSV724 fo
16	14	4	3	18	13	AAQ30310	Oligomer HSV723 fo
17	14	4	3	18	20	AAZ22162	Human c-IAP-1 mRNA
18	14	4	3	22	22	AAQ01590	Human IQGAP2 Cpg I
19	14	4	3	22	22	AAQ01643	Human IQGAP2 5'-UTR
20	14	4	3	26	19	AAV07952	Helicobacter pylori
21	14	4	3	26	19	AAV07952	Helicobacter pylori
22	14	4	3	27	19	AAV07937	Helicobacter pylori
23	14	4	3	29	17	AAV07937	Helicobacter pylori
24	14	4	3	29	17	AAV07937	Helicobacter pylori
25	14	4	3	29	21	AAAS1200	Primer for amplifi
26	14	4	3	31	16	AAV35703	N-terminal primer
27	14	4	3	31	19	AAV67854	Human gene signatu
28	14	4	3	32	20	AAZ27689	Nucleotide fragmen
29	14	4	3	36	18	AAV97603	PCR primer for Ver
30	14	4	3	36	22	AAC30606	Shigella dysenter
31	14	4	3	37	15	AAQ62992	Tomato spotted wil
32	14	4	3	45	22	AAQ55449	Glycophorin antibo
33	14	4	3	45	22	AAQ55450	Oligonucleotide us
34	14	4	3	45	22	AAC88874	Oligonucleotide us
35	14	4	3	45	22	AAC88875	Oligonucleotide TA
36	14	4	3	47	21	AAZ66366	Oligonucleotide TA
37	14	4	3	47	21	AAZ67473	Human map-related
38	14	4	3	47	21	AAZ67533	Human map-related
39	14	4	3	47	21	AAZ67549	Human map-related
40	14	4	3	47	21	AAZ67813	Human map-related
41	13	4	0	50	21	AAV98312	Human MSH6 fragmen
42	13	4	0	15	22	AAV48097	IGFBP3 oligonucleo
43	13	4	0	15	22	AAV48098	IGFBP3 oligonucleo
44	13	4	0	15	22	AAV48099	IGFBP3 oligonucleo
45	13	4	0	16	21	AAV57758	Nucleotide sequenc
	13	4	0	17	16	AAQ92084	Renilla reniformis

ALIGNMENTS

RESULT	1
AAT42655	AAT42655 standard; DNA; 29 BP.
ID	AAT42655 standard; DNA; 29 BP.
XX	
XX	
AC	AAT42655;
DT	
DD	25-FEB-1997 (first entry)
XX	
DE	Primer for amplifying verotoxi
XX	
KW	Verotoxin; Escherichia coli; e
KWK	haemolytic uraemic syndrome; d
XX	
OS	Synthetic.
OS	
NN	WO9630043-A1
PEN	
XX	
PD	03-OCT-1996.

Synthetic.
WO9630043-A1
03-OCT-1996.

25-MAR-1996; 96WO-US04093.
24-MAR-1995; 95US-0410058.
(OPHI-) OPHIDIAN PHARM INC.
Carroll SB, Padhye NV, Stafford DC;
WPI; 1996-505779/50.
Compsn. contg. neutralising antitoxin against E.coli vero-toxin -
used to treat intoxicated individuals, and as a prophylactic against
diarrhoeal disease or extra-intestinal complications of E.coli
infection

PS Example 6; Page 58; 101pp; English.

CC Compositions containing neutralising antitoxin against one or more E.
CC coli verotoxin (VT) can be used to treat intoxicated adults and
CC children with enteric bacterial infections. They may also be used as
CC prophylactics e.g. as a vaccine, against diarrhoeal disease or the
CC development of extra-intestinal complications of E.coli infection,
CC especially haemolytic uremic syndrome. The antitoxin can also be
CC used to detect E. coli VT in a sample. The VT is recombinant,
CC preferably a fusion protein containing a non-VT protein sequence and
CC part of the E.coli VT1 or VT2 sequence. Two primers (AA142655,
CC AA142656) were used to amplify the verotoxin VT-1 A subunit coding
CC sequence and add a histidine tag coding sequence to the subunit
CC sequence. Two primers (AA142655, AA142658) were used to amplify the
CC verotoxin VT-1 A and B subunits and add a histidine tag coding
CC sequence to the subunit sequences.

CC Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.9%; Score 16; DB 17; Length 29;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
DB 9 aaataattatttta 24

RESULT 2

AA151200 standard; DNA: 29 BP.

AA151200;

26-SEP-2000 (first entry)

N-terminal primer for E. coli verotoxin 1 subunit A gene.

VT-1; verotoxin; antitoxin therapy; fusion protein; affinity tag; food;

recombinant production; screening; dairy; anti-bacterial; vaccine;

primer; polyhistidine; ss.

Escherichia coli.

Synthetic.

US6080400-A.

27-JUN-2000.

13-MAR-1997; 97US-0816977.

24-MAR-1995; 95US-0410058.

(OPHT-) OPHIDIAN PHARM INC.

Williams JA, Byrne LM;

WPI: 2000-451195/39.

Bacterial cell for recombinantly expressing bacterial toxins in large
quantities useful for immunization and treatment of bacterial
infections, comprises expression vector encoding bacterial toxin

Example 6; Column 83; 83pp; English.

E. coli verotoxin (VT) type 1 and 2 subunits A and B were cloned into
PET-23b, designed to allow expression of the native proteins containing
C-terminal polyhistidine tags. The VT-1 and VT-2 genes were engineered
to convert the signal sequence methionine codon into a NdeI site to
allow cloning of the amplified genes into the vector without addition of
C-terminal amino acids. The C-terminal primers comprises the
17 codons of each gene fused to the sequence CTCGACC, in order

to add the polyhistidine tag. The primers delete the native stop codons,
and when cloned into PET-23 add a C-terminal extension of Leu-Glu-(His)₆.
VT B chains are small proteins (approximately 8 kDa), so use of a small
affinity tag was preferred (i.e. polyhistidine). A polyhistidine affinity
tag facilitates single step affinity purification of subunits from
periplasmic extracts. However, due to poor recovery of his-tagged VT-1 A
and VT-2 A chains, expression of maltose binding protein (MBP) fused
subunits was undertaken. Due to the toxicity of the VT-2 B subunit,
strict uninduced promoter control is necessary to permit cell viability.
Bacterial host cells expressing a recombinant expression vector encoding
a polyhistidine affinity tag and a portion of the VT-2 B chain are
claimed. The vector is chosen from PET24hisVT2BL+, PET24hisVT2BL- and
PET24hisVT2B, where "L+" indicates that the vector encodes the preprotein
form of the protein and "L-" indicates that the vector encodes the mature
form of the protein. The bacterial cell is capable of expressing large
quantities (40 mg/l) of VT-2B. The toxins are useful for immunizing
non-mammals and for detecting bacterial toxins in environmental samples
including soil, water, industrial samples, biological samples and samples
obtained from food and dairy processing instruments.

Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.9%; Score 16; DB 21; Length 29;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
DB 9 aaataattatttta 24

RESULT 3

AA227689 standard; DNA: 32 BP.

AA227689;

22-DEC-1999 (first entry)

PCR primer for Verotoxin gene.

Verotoxin; VT1; VT2; detection; PCR primer; ss.

Synthetic.

Escherichia coli.

JPI1243996-A.

14-SEP-1999.

27-FEB-1998; 98JP-0047677.

27-FEB-1998; 98JP-0047677.

(TOYM) TOYBO KK.

WPI: 1999-603716/52.

An oligonucleotide for amplification of verotoxin - useful in the
detection of inactivated verotoxin gene by transfer of a foreign DNA
fragment

Claim 11; Page 9; 10pp; Japanese.

This sequence represents a PCR primer of the invention. The primer is
used for amplification of the E. coli verotoxin (VT) gene. The
oligonucleotide is useful for detection of inactivated VT gene by
transfer of a foreign DNA fragment. Simple, rapid and specific
amplification of VT gene from environmental factors is achieved using the
oligonucleotide of the invention.

Sequence 32 BP; 12 A; 2 C; 4 G; 14 T; 0 other;

```

FT FT /note= "N-methyl-8-oxo-2'-deoxyadenine"  

FT FT 15  

FT FT /*tag= m  

FT FT /mod_base= OTHER  

FT FT /note= "N-methyl-8-oxo-2'-deoxyadenine"  

FT FT 17  

FT FT /mod_base= n  

FT FT /*tag= n  

FT FT /mod_base= OTHER  

FT FT /note= "N-methyl-8-oxo-2'-deoxyadenine"  

PN PN W09118997-A.  

XX XX  

PD PD 12-DEC-1991.  

XX XX  

PF PF 24-MAY-1991; 91WO-1003680.  

XX XX  

PR PR 14-JAN-1991; 91US-0640554.  

PR PR 25-MAY-1990; 90US-0529346.  

XX XX  

PA (GILE-) GILEAD SCIE INC.  

PI Matteucci MD, Krawczyk S;  

XX WPI; 1992-007480/01.  

DR WPI; 1992-007480/01.  

XX  

PT New sequence-specific non-photo-activated crosslinking a  

PT bind to the major groove of duplex DNA and are esp. usef  

PT treating latent infections e.g. HIV  

XX  

PS Example 4; Page 29; 42pp; English.  

XX  

CC This oligomer contains an inverted polarity region forme  

CC o-xylosa dimer synthon. Residues 11 and 12 are linked vi  

CC o-xylosa group (i.e. nucleotides that have xylose sugar  

CC the o-xylene ring). The sequence is designed to target th  

CC Simplex virus I beginning at nucleotide 10996 and to cov  

CC cross-link to it. See also AAQ20151-Q20160.  

XX  

SQ Sequence 18 BP; 12 A; 1 C; 0 G; 5 T; 0 other;  

  

Query Match 4.3%; Score 14; DB 13; Length 11  

Best Local Similarity 100.0%; Pred. No. 6.9e+03;  

Matches 14; Conservative 0; Mismatches 0; Indels  

  

QY 159 aaatataaaataaa 172  

| | | | | | | | | |  

Db 2 aaatataaaataaa 15  

  

RESULT...14  

AAQ20160  

ID AAQ20160 standard; DNA; 18 BP.  

XX  

XX AAQ20160;  

XX  

DT 01-APR-1992 (first entry)  

DE Cross-linking oligomer 723 to target Herpes Simplex Virus  

XX deoxyribonucleic acid; major groove; HSV;  

KW inverted polarity region; covalent cross-linking group s  

KW Synthetic.  

OS  

FH Key Location/Qualifiers  

XX modified_base 1 /*tag= a  

FT FT /mod_base= OTHER  

FT FT /note= "N-methyl-8-oxo-2'-deoxyadenine"  

FT FT 2  

FT FT /*tag= b  

FT FT /mod_base= OTHER

```